POTASSIUM MOVEMENT IN LAKELAND FINE SAND AND EFFECTS ON YIELD AND COMPOSITION OF FLUE-CURED TOBACCO . (Nicotiana tabacum L.)

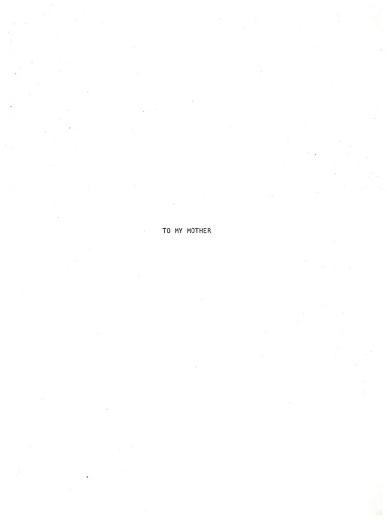
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Abstract of Dissertation Presented to the Graduate Council of the University of Florida in Partial Fulfillment of the Requirements for the Degree of Doctor of Philosophy

POTASSIUM MOVEMENT IN LAKELAND FINE SAND AND EFFECTS
ON YIELD AND COMPOSITION OF FLUE-CURED TOBACCO
(Nicotiana tabacum L.)

By

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Chairman: William K. Robertson Major Department: Soil Science

Potassium (K) movement in Lakeland fine sand was studied in the field with and without indicator crop, and in the laboratory using reconstituted soil columns. To determine effects of K movement on yield and composition of flue-cured tobacco (Nicotiana tabacum L.), different slowly soluble and soluble fertilizers containing K were tested. Although there were observed differences in yields of flue-cured tobacco and in K, total alkaloid, and total reducing sugar contents of the tobacco leaves from two years' field experiments, the differences did not indicate any advantage of the slowly soluble K fertilizers over the tobacco-grade fertilizer applied split. Results from field and laboratory experiments showed extent of K movement as affected by amount of water applied to the soil. Results from miscible displacement studies conducted in the laboratory supported the observation from both field and laboratory experiments that K was lagging behind the water moving downward.

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INTRODUCTION

Common sources of potassium (K) applied to agricultural crops are very soluble in water. Their use, particularly at high rates in combination with other fertilizers, have fostered problems associated with high salt concentration in the plant-root environment. In sandy soils, during periods of heavy rainfall, leaching may reduce K concentration to a level of deficiency. To alleviate high salt concentration and to minimize leaching losses, split applications of the fertilizer K are recommended and additional applications are suggested after unusually heavy rains. Such is the case for flue-cured tobacco (Nicotiana tabacum L.) production in Florida. Slowly soluble sources of K, which are less susceptible to leaching, are now available commercially and others are currently undergoing testing for their effectiveness. More products will probably become available commercially in the near future. These sources may prove to be effective in satisfying the crop's requirement for K, possibly in only one application and at a relatively low rate.

Flue-cured tobacco is an important cash crop in Florida. The market value of the crop depends not only on the total yield but also on the grade of the cured tobacco leaves as determined by federal tobacco inspectors. The tobacco grade is an index of the quality of cured leaves which is related to the chemical composition. Yield and quality are markedly influenced by climate and cultural practices. Soil fertilizer management is therefore a very important aspect of the cultural practices in commercial tobacco farms and tobacco growers are very much aware of this. For example, farmers apply excessive amounts of fertilizers to

ensure the crop of an available nutrient supply under conditions of high rainfall. If excess fertilizer is not leached by rainfall it may be leached by irrigation.

There has been much work done on nutrient movement in the soil under field, laboratory, and greenhouse conditions. Nitrogen (N) and phosphorus (P) have received most attention because of their importance in environmental pollution. Movement of K is of more concern relative to its availability to plants since it is not considered a pollution hazard. Studies of K movement in the soil are more valuable if carried out under field conditions with a growing crop since it allows evaluation of factors affecting K movement in relation to plant uptake and other growth parameters. Since tobacco takes up K in large quantities, it is a good indicator crop to study K movement.

The objectives of this research were to study the effect of soluble and slowly soluble K fertilizers on yield, composition, and quality of flue-cured tobacco and to determine the effect of rainfall and irrigation on the movement of K from these fertilizers in Lakeland fine sand. These objectives were determined in the field and laboratory.

LITERATURE REVIEW

Chemically, flue-cured tobacco consists mainly of organic substances such as alkaloids, organic acids and bases, carbohydrates, essential oils, resins, and other nitrogenous constituents (Hawks, 1970). Various analyses are made to predict quality of flue-cured tobacco. The most common analyses are: total N, protein N, ∞ -amino N, nicotine, reducing sugars, starch, non-volatile acids, and petroleum ether extractables. The quantity as well as the balance of these constituents determine the quality of tobacco (Hawks, 1970).

The flavor and pleasing quality of the taste of smoke is directly related to the content of the nitrogenous constituents (Hawks, 1970).

A high total N content produces a strong or more pungent taste. If the content of nitrogenous constituents is low, the taste is flat and unpleasing. High content of oxalates and insoluble carbohydrates impart a bitter and irritating taste to the smoke. The lower the «-amino N content of tobacco the more aromatic and pleasing the smoke will be.

A good quality tobacco is also associated with a high sugar content if the N balance is maintained. Sugar content of the leaf is negatively correlated with N content (Woltz, Reid, and Colwell; 1948).

A high K content of flue-cured tobacco has been used as a measure of quality (Hawks, 1970). The quality factors related to K content are concerned with the physical appearance which determines the grade of the tobacco leaves in the market. A high K content is believed to produce deeper orange color of leaves. Hawks, Terry, and Collins (1973) observed no significant response of yield and quality of flue-cured tobacco to

rates of K fertilization in some North Carolina soils.

Nelson and McEvoy (1945) as cited by McCants and Woltz (1967) obtained evidences suggesting that tobacco plants can build up a reserve of K in the early stages of growth sufficient to sustain the later phase of development. Deficiency symptoms occurred when K supply was eliminated four weeks prior to harvesting. On the other hand, when the K supply was maintained until two weeks prior to harvesting, the reserve in the plant sustained the remaining development without additional absorption.

The availability of K from some slowly soluble K fertilizers to flue-cured tobacco was studied in a field experiment by San Valentin, Robertson, and Lundy (1971). The results showed that lower rates of Osmocote 12-0-34 (12% N, 0% P, 34% K) and sulfur-coated $\rm K_2 SO_{l_4}$ (SCK) produced yields and value of flue-cured tobacco that were not significantly different from those using higher rates of commercial 4-3.5-10 and granular $\rm K_2 SO_{l_4}$, which are both soluble sources of K. The total K uptake in the harvested leaves followed the order: Osmocote 12-0-34 > commercial 4-3.5-10 (applied split) > SCK (26% coating) > SCK (21% coating) > $\rm K_2 SO_{l_4}$. MgSO_{l_4} (K-Mag) > granular $\rm K_2 SO_{l_4}$.

The role of K in the plant is not clearly understood unlike other nutrients because studied on K are confounded with difficulties associated with its being soluble in the cell sap and not being synthesized with any specific compound in the plant. A review of literature has been written on the role of K in photosynthesis and respiration by Jackson and Volk (1968), on carbohydrate metabolism and translocation by Liebhardt (1968), on effects of K on organic acid and nonprotein N content by Teel (1968), and on effects of K on enzymes by Wilson and Evans (1968). One of the most recent findings on the role of K in photosynthesis is on the closing

and opening of the stomata which has been found to be dependent on the movement of K in and out of the guard cells (Jackson and Volk, 1968).

On carbohydrate metabolism, K has been observed to reduce condensation of simple carbohydrates to more complex forms (Liebhardt. 1968). Numerous enzyme systems in the plant require K in their reactions and many of these enzymes are involved in protein or N metabolism (Wilson and Evans, 1968).

The K requirement of crops varies (Fried and Broeshart, 1967).

Tubers and root crops, tobacco, pineapple, and sugar cane remove large amounts of K usually more than 200 kg K/ha. Vegetables such as tomatoes, cauliflower, and cabbage also remove large amounts of K from soil. Grain crops and legumes remove less, the former from 67 to 78 kg K/ha and the latter from 45 to 123 kg K/ha.

Nutrient movement from the soil to the plant may be described by three mechanisms: 1) movement through the soil to the root surface,

2) movement from the root surface to the interior of the root, and

3) movement from root to various parts of the plant. Nutrients may reach the root surface by root interception, mass-flow, and diffusion (Barber, 1962 and Barber, Walker, and Vasey, 1963). Movement by mass-flow is related to water movement along the transpiration stream. As soil water around the root becomes depleted, a hydraulic gradient develops causing water movement from regions of greater water content to regions immediately surrounding the roots. The significance of mass-flow depends on the volume of water absorbed by the plant and the concentration of the nutrient in the soil solution. Movement by diffusion is over much shorter distances than mass-flow (Drew, Vaidyanathan, and Nye; 1967).

Among the factors limiting nutrient uptake by plant roots are the amount and distribution of roots in the soil. Roots come in contact

with the nutrients as they grow and the maximum amount of nutrients which can be contacted depends on the volume of roots formed and the concentration of the nutrient in that particular volume of soil where the roots are growing (Barber, Walker, and Vasey; 1963).

Howard and Adams (1965) showed that cotton root extension in the soil is related to the ratio of Ca to total cations. When Ca content was less than 20% of total cations, root growth decreased sharply.

Adams and Lund (1966) showed restriction of primary root growth by Al in three acid soils of widely different properties. A relationship was observed between Al concentration of soil solution and root growth but not with exchangeable Al.

The amount of K absorbed by roots is determined by concentration or more appropriately the activity of K in the bulk soil solution which in turn is dependent on the electrochemical potential of K and other ions surrounding the exchange complex forming the diffuse double layer. The electrochemical potential of ions in the exchange phase can not be measured, nor it is directly proportional to the concentration of ions in the equilibrium soil solution (Beckett, 1967). This problem is avoided by measuring the differences between the electrochemical potential of two ions which is constant in all parts of the solution, double layer, or solid phase at equilibrium (Beckett, 1972). At equilibrium, the activity ratio of K to Ca in soil solution, a_{K}/a_{Ca} , is proportional to their difference in electrochemical potential. The similarity in behavior of Ca and Mg made it possible to express a(Ca+Mg). Measurement of this ratio has been considered a useful indication of K supply to plants (Beckett, 1964; Wild, Rowell, and Ogunfowora, 1969; and Graham and Fox, 1971). Measurement of this ratio in relation to the amount of

labile K is known as quantity-intensity (Q/I) technique as proposed by Beckett (1967).

Lime may convert exchangeable K to nonexchangeable forms and induce adsorption of K by soil colloids (Peech and Bradfield, 1943). On four North Carolina soils made homoionic with respect to Ca and Al. Thomas and Coleman (1959) observed that Ca-soil retained more K than did Al-soil due to an increase in effective cation exchange capacity (CEC) with Ca saturation.

Murdock and Rich (1965) studied the effect of lime on K availability in Nason silt loam. Potassium fixation occurred at high pH but did not appear to be associated with greater collapse of the vermiculite present in the soil. It was thought to be due to less competition by ${\rm H_30}^+$ for the specific adsorption sites for K.

Gammon (1957) reported that the first effect of liming on soil K was to reduce exchangeable K. A reduction from 26 to 8 ppm of exchangeable K in Leon fine sand was observed one year after application of 5.9 tons of lime per hectare which changed the pH of the soil from 4.6 to 5.9. Repeated K fertilization following liming increased the exchangeable K in the soil over a three-year period. Bartlett and McIntosh (1969) observed K deficiency in tomato seedlings on a spodosol Ap which was high in extractable Al and pH-dependent CEC, after liming with ${\rm CaCO_3}$, ${\rm CaSiO_3}$, ${\rm CaHPO_4}$, and ${\rm CaF_2}$ to pH 6.3. It was hypothesized that positive charges associated with Al were probably neutralized by ${\rm OH^-}$, ${\rm PO_4}^{-3}$, or ${\rm F^-}$ thereby increasing the CEC but decreasing percentage saturation of the exchange complex by K.

Coating soluble K fertilizers reduce K losses by leaching and increase K availability to plants (Lawton, 1961). Various materials

have been used experimentally as coatings, the most important being waxes, polymers, and sulfur (Hauck and Koshino, 1971). Examples of polymers used are: acrylic resins, polyurethane, polystyrene, polyethylene, vinyl acetate, polyvinyl chloride-polyvinylidene chloride, and polyacrylate nitrile.

Uncoated slowly soluble K fertilizers such as potassium polyphosphate (KPP) and potassium-calcium pyrophosphate (KCP) have also been studied (Chandler and Musgrave, 1944; Lutrick, 1958; DeMent and Stanford, 1959; Caldwell and Kline, 1963; Hagin, 1966; and Engelstad, 1968). Their studies indicated that those slowly soluble K fertilizers are as effective as the soluble fertilizers and they are less likely to leach.

The efficiency of fertilizers also varies with particle size of the material. Hagin (1966) observed that a powdered form of a KPP was more efficient than the granulated form in calcareous and alkaline soils.

DeMent, Tenman, and Bradford (1963) observed that K uptake by corn decreased with increasing particle size of KPP mixed through the soil while the reverse was observed for surface applied KCP.

The retention of KC1, K_2SO_{4} , and K_3PO_{4} were investigated by Ayres and Hagihara (1953) on some tropical soils of Hawaii. The order of retention of K was as follows: $K_3PO_{4} > K_2SO_{4} > KC1$. Greater retention of K was observed from the application of a mixture of ammophos and KC1 particularly with higher ratio of phosphate to K. Superphosphate or treble superphosphate application with KC1 did not significantly reduce K losses possibly because Ca and H were strongly held in contrast with NH $_4$ from ammophos. The less strongly held NH $_4$ could allow greater retention of K when ammophos was applied with KC1. The observed increase in retention by adding phosphate was later clarified by the findings of

Mekaru and Uehara (1972). They observed that the CEC of a Hydrandept increased from 28 to 127 me/100 g by addition of phosphate. They explained that specifically adsorbed anions such as phosphate render the surface of constant potential colloids more negative by changing the zero point of charge to a lower pH.

The efficiency of soluble K fertilizers also varies according to the method of placement in the field. Volk and Bell (1945) observed greater K losses by band placement of fertilizer than by broadcast in Norfolk fine sand. Nolan and Pritchett (1960) also observed greater losses of K from KCl by band placement than broadcast in Arredondo fine sand with sweet potato as an indicator crop. Adams et al. (1967) observed more efficient utilization of K from KCl by Coastal bermudagrass when 154 kg/ha of K was applied in four equal applications than when applied only once.

Leaching of K is influenced by the capacity of the soil to retain cations. This property is determined by the amount and type of clay minerals present and organic matter content of the soil. Mica, montmorillonite, and vermiculite are the clay minerals contributing to the CEC of most soils. The adsorption of K by these minerals may sometimes lead to a condition where K is rendered unavailable or difficultly available to plants. The process is known as fixation although the term "fixed" K is regarded as ambiguous since this fraction of soil K depends on the extraction procedure.

Stanford (1947) showed that micaceous minerals are responsible for K fixation under moist soil conditions while montmorillonite did not fix K except on drying. Drying increased the amount of K fixed by illite but under moist conditions acidified illite fixed little K.

Studies by Mortland, Lawton, and Uehara (1957) indicated that K

fixed by vermiculite and montmorillonite is more easily released by illite and muscovite as shown electrodialysis, chemical extraction and plant growth procedures.

While most studies on K fixation were carried out on crystalline minerals, K fixation by amorphous minerals has received considerable attention. The information on K fixation by amorphous aluminosilicates in soils is scarce because of the difficulty in the extraction of amorphous aluminosilicates. Results of a study by van Reeuwijk and de Villiers (1968) on synthetic amorphous aluminosilica gels showed K fixation in varying amounts measured against exchange by different cations.

In Florida, the clay fraction generally constitutes a very small fraction of the whole soil. Hence, organic matter plays a dominant role in the soil's capacity to retain cations (Fiskell, 1970). Florida soils were found to possess a great portion of pH-dependent CEC (Fiskell and Zelazny, 1971 and Zelazny and Fiskell, 1971). Spencer (1954) studied Lakeland, Blanton, Ft. Meade, and Eustis series and found that colloids of these sandy soils adsorb K very weakly and this accounts for the rapid leachability of K in these soils.

The mineralogical composition of the clay fraction of some Florida soils has been studied by Fiskell and McCaleb (1953) and Fiskell et al. (1958). The dominant clay minerals they found in Florida soils were kaolinite and gibbsite. Montmorillonite, vermiculite, quartz, and feldspars were also found but in relatively small amounts.

The contribution of organic matter to soil CEC based on Ba and K adsorption at various pH values was studied by McLean and Owen (1969).

Their results indicated that organic exchange sites active for Ba adsorp-

tion from buffered ${\rm BaCl}_2$ solution at low pH were not active for K adsorption from unbuffered KCl solution of similar pH. This observation, they explained, was due to nonexchangeable Al associated with organic matter.

Water movement is a very important factor determining extent of K leaching in soils. Water movement takes place through interconnected pores in the soil. Under saturated conditions, the pores are filled with water but the large pores are more effective as conducting channels for water flow. As the soil becomes less saturated, the small pores become more important as conducting channels but the flow path becomes more tortuous (Baver, Gardner, and Gardner; 1972). Thus, the ability of the soil to transmit water or hydraulic conductivity, $K(\theta)$, decreases as water content decreases (Miller and Klute, 1967). Although the saturated hydraulic conductivity is greater in sandy soils than fine textured soils, the change in conductivity with water content is less abrupt in fine textured soils because of wider pore size distribution.

The distribution of pore size and their shapes results in a distribution of velocities of water flow through the interconnected pores.

The distribution of flow velocities as well as molecular diffusion are the two primary processes contributing to mixing of fluids during miscible displacement (Day, 1956 and Biggar and Nielsen, 1967).

Adsorption, desorption, and precipitation of solutes may take place as solutes move along with water during flow through the soil. Ion exchange therefore is a very important consideration in modelling for solute movement through the soil. Boast (1973) presented a review of models to describe solute movement in the soil, specifically those based on classical mathematical macroscopic continuum theories. Other models are based upon statistical theories (Day, 1956 and Nakano and Murrmann,

1971) and finite difference theories. Lai and Jurinak (1972) applied the finite difference method to the solution of the material balance equation to describe Na adsorption in one-dimensional flow through the soil. Sodium adsorption on originally Ca-saturated Yolo soil agreed with the values obtained by numerical solution of the theoretical equation.

Rible and Davis (1955) applied the statistical theory together with chromatographic theory of ion exchange in soil columns. The assumptions required for the theory are as follows: I) occurence of instantaneous ion-exchange reactions so that the solution flowing through the exchange column is considered in equilibrium with the exchanger, 2) there are negligible effects of ionic diffusion, channeling, turbulent flow, and alteration of the void ratio during solution flow, 3) the exchange capacity of the exchanger is unaltered during solution flow, and 4) the exchange constant remains unchanged over the range of concentration involved. Assumptions I and 2 are not recognized to be realistically applicable to soil columns. Their data from Yolo fine sandy loam soil indicated good agreement between theory and experimental results for the replacement of Ca by Mg and by Na. Fair agreement was obtained when Ca replaced Na but not when Ca replaced Mg.

Thomas and Coleman (1959) applied a chromatographic theory to leaching of fertilizer salts in soils. The chromatographic theory employed was one by Walter which considered CEC and ion-exchange equilibrium constant as the only two factors involved. The theory agreed with values of K distribution in the soil initially made homoionic with respect to Ca and Al and eluted with a solution of KCl and then followed with water. With Na as the initial saturating cation, the theory failed

to predict K distribution unless K added was large relative to the amount of Na in the column. Hydrolysis and change in effective CEC of the soil were speculated to be the possible reasons for the lack of agreement with the Na system.

Simultaneous flow of water and solutes under saturated or unsaturated soil water conditions have been studied in the past. In most cases studies have been conducted using soil columns in the laboratory. It has been necessary to apply several assumptions and complex boundary conditions for the calculations of the solutions to the differential equations describing simultaneous flow of solutes and water. For example, most work done in the past assumes a constant flow velocity and a homogeneous porous medium. In the study by Bresler and Hanks (1969) a method was devised for estimating the distribution of a non-interacting solute in the soil in response to varied wetting and drying conditions and allows for a time-dependent flow velocity. Their equation considered only the effect of viscous flow since the effect of diffusion on the flow of solutes was assumed to be very small compared to that of viscous flow.

Warrick, Biggar, and Nielsen (1971) studied simultaneous movement of water and solute for an unsaturated soil under field conditions. The field data indicated that the distribution of chloride (Cl⁻) in the soil after it was first applied with irrigation water and then followed with Cl⁻-free water can be predicted quantitatively by combining an equation of solute movement and water movement through an unsaturated soil.

Miller, Biggar, and Nielsen (1965) studied Cl⁻ movement in relation to water movement in Panoche clay loam free of vegetation. Three methods of water application were compared, 1) continuous ponding, 2) intermittent ponding with 15 cm of water, and 3) intermittent ponding with 5 cm of water. Intermittent leaching and smaller amount of leaching water moved the salt more like a slug, characterized by prominent peaks whereas under continuous ponding, the salt distribution was more uniformly spread through the soil. The experiment showed that C1⁻ movement may be altered with the method of water application.

MATERIALS AND METHODS

Field Experiments

Field Experiments on Flue-Cured Tobacco

Field experiments were conducted at the Agricultural Research Center at Live Oak, Florida in 1972 and 1973 to determine the effects of different K fertilizers on yield, composition and quality of flue-cured tobacco (Nicotiana tabacum L.). The various K fertilizers studied are summarized in Tables 1 and 2 for the 1972 and 1973 experiments, respectively, together with a description of the treatments associated with each fertilizer. The soil in the area is a Typic Quartzipsamment, siliceous, thermic, and coated, known commonly as Lakeland fine sand.

Both 1972 and 1973 experiments had treatments that were factorial arrangements of fertilizers or method of application and rates of application replicated once in three blocks. However, in the 1972 experiment there were two additional treatments (Table 1). The experimental plots contained four rows 12 m long and 1.2 m apart.

In these experiments the tobacco-grade fertilizer was used as a control. It was banded along the rows as recommended, half at planting and half two weeks after planting. The bands were about 8 cm wide and about 10 cm from the plant row. Bands of fertilizer were covered by

Table 1. Treatments in 1972 field experiment on flue-cured tobacco $\overline{1}^{\prime}$

, ment						
ė.	Į.	Materials			Method of application	Comments
_	NH4N03-ordinary	$^{\mathrm{NH}_{4}\mathrm{NO}_{3}} ext{-ordinary superphosphate-}^{\mathrm{K}_{2}\mathrm{SO}_{4} ext{-MgSO}_{4}}$	K ₂ S04-	MgS04	½ at planting, ½ two wk after	Tobacco-grade fertilizer
2	=	Ξ	Ξ	=	All at planting	=======================================
~	=	=	=	=	All K at planting, 늘 of base N and P at planting and 글 two wk after planting	K source was sulfurcoated $K_2SO_{\rm L}$ with 36% coating $(SCK,~36\%)\underline{2}/$
4	=	Ξ	Ξ	=	All K at planting, $\frac{1}{2}$ of base N and P at planting and $\frac{1}{2}$ two wk after planting	K source was sulfur- coated $K_2SO_{\rm L}$ with 18% coating $(5{\rm CK},~18\%) \underline{2}/$
2	Osmocote (7-6-17.5)-MgSO $_{\mu}$	17.5)-MgSO ₄			All at planting	Osmocote (7-6-17.5) is resin-coated 7-6-17.5
9	Osmocote N-ordi	Osmocote N-ordinary superphosphate- ${\rm K_2S0_{4^-}M950_{4}}^{11}$	ate-K ₂	SO4-MgS	· · · · · · · · · · · · · · · · · · ·	N source was resin-coated $\mathrm{NH}_4\mathrm{NO}_3$, K was $\mathrm{K_2SO}_4$
7	NH ₄ NO ₃ -ordinary	$^{\mathrm{NH}_{4}\mathrm{NO}_{3}} ext{-ordinary superphosphate-MgSO}_{4}$	4080		글 at planting, 눌 two wk after	No K applied
œ	0smocote (7-6-17.5)-MgSO ₄	17.5)-MgS0 ₄			All at planting	Similar to middle rate of treatment 5 except all N including sidedressing, applied at planting

 1 /Treatments 1 to 6 applied at rates equivalent to 450, 900, and 1800 kg/ha of tobacco-grade fertilizer (4% N, 3.5% P, 10% K, and 1.29% Mg) and all treatments except 8 received 34 kg N/ha as NH $_{4}$ NO $_{3}$ five wheeks a ferr planting, banded beside the row.

arter planting, banded beside the row. 2^{\prime} Materials provided by Tennessee Valley Authority.

Table 2. Treatments in 1973 field experiment on flue-cured tobacco $\frac{1}{1}$

Treat- ment No.	t- Materials	Method of application	Comments
-	$^{ m NH}_4 ^{ m NO}_3$ -ordinary superphosphate- $^{ m K}_2 ^{ m SO}_4$ - $^{ m HgSO}_4$ $^{\frac{1}{2}}$ at planting, $^{\frac{1}{2}}$ two wk after	½ at planting, ½ two wk after	Tobacco-grade fertilizer
2	Swift (0-6.5-25)-IBDU-MgSO $_{ m t}$	All at planting	Swift (0-6.5-25) and IBDU are slowly soluble materials
~	Swift (0-6.5-25)-WH ₄ NO ₃ -MgSO ₄	= = =	Same as treatment 2 except N source was NH $_{4}^{\rm NO}_{3}$
4	$^{\mathrm{NH}_{4}\mathrm{NO}_{3}}$ -ordinary superphosphate-K $_{2}$ SO $_{4}$ -MgSO $_{4}$	All K at planting, ½ of base N and P at planting and ½ two wk after planting	K source was sulfur- coated K_2S_0 , with 26% coating $^2(S^4_0K, 26\%)$ $2/$
5	0smocote (7-6-17.5)-MgS0 $_{ m t}$	All at planting	Osmocote (7-6-17.5) is resin-coated 7-6-17.5
9	$^{ m NH}_{\rm L}{ m NO}_3$ -ordinary superphosphate-MgSO $_{ m L}$	½ at planting, ½ two wk after	No K applied

L/Treatments 1 to 6 applied at rates equivalent to 450, 900, and 1800 kg/ha of tobacco-grade fertilizer (4% N, 3.5% P, 10% K, and 1.29% Mg) and all treatments received 34 kg N/ha as ${\rm NH_4M0_3}$ five weeks after 2/haterial provided by Tennessee Valley Authority.

plowing along the rows. The tobacco-grade fertilizer was applied at three rates: 450, 900, and 1800 kg/ha. Regardless of fertilizer rate, the N was sidedressed at 34 kg N/ha as $\mathrm{NH_4NO_3}$ five weeks after planting. The experimental plots were irrigated with approximately 2.5 cm of water when plants started to wilt.

Other practices recommended in flue-cured tobacco production were also employed in the experiments. These included weed control, pest control, topping, chemical sucker control, harvesting or priming of mature leaves, and curing of harvested leaves (Hawks, 1970). The harvested leaves were cured in gas-fired bulk barns at 35 C to 49 C during the first two weeks and 49 C to 77 C during the next three to four weeks.

Cured leaves from the two inner rows of each plot were weighed.

The leaves were graded by federal tobacco inspectors. Yields are reported in kilograms per hectare and their values in dollars per hectare, calculated using the season's average values for each grade of flue-cured tobacco (type 14) reported by U. S. D. A. Consumer and Marketing Service.

Samples for plant tissue analyses were obtained from the bulk samples of cured leaves representing each of the five primings from each plot and oven-cried at 70 C. The oven-dried leaves were crushed by hand, mixed, and midribs were removed. Samples of midrib-free tobacco leaves were ground to pass through a 20-mesh screen in a Wiley mill and stored in air-tight plastic containers.

One-gram samples of the ground leaves were placed in 50-ml beakers and ashed in a muffle furnace at 450 C. After cooling, the ash was moistened with distilled water, dissolved in 20 ml of 40% HCl, and

evaporated to dryness on a hot plate. About 10 ml of distilled water and another 2.25 ml of 40% HCl solution were added and the resultant solution was boiled for a few minutes without drying. The solution was then filtered through #40 Whatman filter paper and diluted to volume in a 50-ml volumetric flask. Potassium was determined using Beckman model B flame spectrophotometer and Ca and Mg using Perkin-Elmer model 303 atomic absorption spectrophotometer. The amounts of K, Ca, and Mg in the tissue are reported as percent of the oven-dry sample.

The analyses of total alkaloids and total reducing sugars were performed in the laboratory of Dr. W. W. Weeks at North Carolina State University.

Soil samples from 0-15, 15-30, 30-45, and 45-60 cm depth were collected weekly from selected plots starting two weeks after planting to determine movement of K. Samples were taken midway between plants in each of the two middle rows using a sampling tube 2.54 cm in diameter. Samples from the two holes were composited, air-dried, and passed through a 2-mm sieve. Five-gram subsamples were shaken with 25 ml of $\frac{N}{4}$ NH $_4$ OAC (pH 4.8) and the extracts filtered through #40 Whatman filter paper prior to analysis for K.

Analyses of variance for yield, value, and soil and plant tissue analyses data from both experiments were performed by Dr. F. G. Martin of the University of Florida Statistics Department using an IBM 360 computer. The 1972 experiment was not a balanced design, therefore additional analyses were required for the two additional treatments. The additional analyses consisted of the comparisons between effects of treatment 7 and those of treatments 1 to 6 and between effects of treatment 8 and that of the second rate of treatment 5 (Table 1).

Duncan's multiple range tests (DMRT) were also performed where effects were statistically significant. Where rate of fertilizer application effects were significant, regression equations, either linear, quadratic, cubic, or quartic, were calculated. Analyses of variance were performed both on overall priming and individual priming data.

Field Experiments on Potassium Leaching

The two field experiments described were almost an acre in size and required tractors and other equipment for operation. Use of those equipment made it difficult to set up tensiometers and suction equipment for a sophisticated leaching study with a growing crop. Therefore, a separate field experiment was set up without tobacco plants to study K movement in relation to movement of water in the soil.

The design of the K leaching experiment was similar to that of Warrick et al. (1971) using tensiometers and soil solution samplers. The tensiometers and soil solution samplers were installed at random location within a 3 m by 3 m area at depths of 15, 30, 45, 60, 75, and 90 cm below the surface. Two sets of soil solution samplers and two sets of tensiometers were used. Each sampler was connected to a suction flask and the flasks were connected in series to a vacuum pump. The method of connecting each of the soil solution samplers to the suction flask is shown diagramatically in Figure 1. This set up permitted quick and independent suction and withdrawal of a small volume of soil solution from each sampler. Soil water tensions were determined through the mercury manometer of each tensiometer.

A solution sampler consisted of a porous ceramic sup 1.12 cm outside diameter, attached to one end of a plastic tube similar to that used for

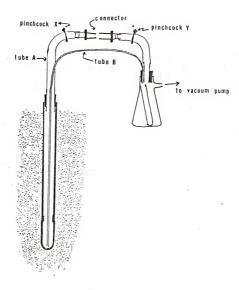


Figure 1. Soil solution sampling apparatus used in the field experiment on $\ensuremath{\mathrm{K}}$ leaching.

the tensiometers. The other end was connected with two Tygon tubes (tubes A and B in Figure 1) through a rubber stopper. The large tube (A) was 0.63 cm in diameter and served as the main suction line. The small tube (B) was 0.16 cm in diameter and delivered solution at the bottom of the sampler to the test tube in the suction flask. The small tube also provided for flushing of the samplers with distilled water prior to each soil solution sampling.

After a few preliminary runs with the set up, a procedure for quick solution sampling was developed wherein small volume increments of soil solution could be extracted and allowed to accumulate. This was important because of the time-dependent movement of water and solutes in the soil. The procedure consisted of applying suction for 3 to 5 min and closing tube A at point X and Y (Figure 1) with pinchcocks, then detaching the tube at the connector. Releasing slowly the pinchcock at point X allows air to enter the solution sampler which would force the solution that was present in the ceramic cup to the test tube in the suction flask. When solution flow stopped as determined visually, tube A was again attached to the vacuum line and pinchcocks at points X and Y were removed. This procedure was repeated several times to all samplers until about 5 ml of soil solution was obtained from each sampler.

A water sprinkler was calibrated to determine output and uniformity of water distribution by using several liter cans placed at various distances from the sprinkler. Three one-hour tests showed that output was 2.5 cm/hr and uniform throughout the experimental area.

The first of two experiments was undertaken on July 3, 1973. Soluble fertilizer (7-6-17.5) was spread evenly over an area of 3 m by 3 m at a rate to supply 180 kg K/ha. About 3 cm of the surface soil was

cultivated with a hand trowel to mix the fertilizer in the soil. After setting the tensiometers and flushing all soil solution samplers, the sprinkler was operated for 2 hr providing a total of 5 cm of water.

Tensiometer readings were recorded before application of water and at 1, 7.5, 22, 74, and 122 hr after application of water. Soil solution samples were collected before application of water and after 1, 7.5, 22, 74, and 122 hr. On July 8, 1973 another 2.5 cm of water was applied and solution samples were collected on July 9 and July 13, 1973. There were rainfalls of 1.65 and 2.03 cm during this period, on July 9 (four hours before sampling) and July 12, respectively.

The second experiment was initiated on October 18, 1973. A 3 m by 3 m area was divided into two subplots. In each, one set of tensiometers and one set of soil solution samplers were installed at random location. Resin-coated 7-6-17.5 was applied to one plot and uncoated 7-6-17.5 to the other. Each fertilizer was applied at a rate to supply 180 kg K/ha. The top 3 cm of the soil was mixed with the fertilizer as in the previous experiment and then water was applied continuously for 6 hr at 2.5 cm/hr. Tensiometer readings were recorded before, at the end of water application, and 96 hr after water application. Soil solution samples were collected at 0 and 96 hr after application of water. All soil solution samplers were flushed with distilled water prior to each sampling.

The soil solution samples were analyzed for K. Samples from the second experiment were also analyzed for Ca and Mg in addition to K.

Laboratory Experiments

Leaching Experiments

Laboratory leaching experiments were carried out on reconstituted soil columns to study K leaching under conditions of limited evaporation of water from the soil surface. The water table was controlled and the system did not include plants.

A cylindrical polyvinyl chloride tube, 10 cm inside diameter, 142 cm long, 0.63 cm thick with a lucite sheet glued to the bottom, was used in these experiments. The bottom contained a hole bored in the center into which a tube connector was glued and a piece of Tygon tubing attached. The Tygon tubing was provided for drainage and for regulating the depth of the water table. Ten tensiometer-samplers were installed at 10 cm interval along the side of the cylinder.

Bulk soil samples representing 0-10, 10-20, 20-30, 30-40. 40-50, 50-60, 60-70, 70-80, 80-90, and 90+ cm depths were collected. The samples were air-dried, passed through a 2-mm sieve and stored in plastic bags. Based on bulk densities determined from undisturbed soil cores, adequate amounts of each sample were weighed and placed into the cylinder to reconstitute the soil profile. To minimize particle segregation during packing, a large funnel attached to a rubber tube 2 cm in diameter and about 150 cm long was used to deliver soil to the bottom of the plastic cylinder. With the end of the rubber tube touching the bottom of the plastic cylinder, the appropriate soil was poured slowly into the funnel. While slowly raising the tube to allow the soil to flow out, the funnel was continually filled with the appropriate soil.

Two reconstituted columns of Lakeland fine sand were saturated with distilled water from the bottom of the columns and after a week under saturated condition, the columns were allowed to drain to a water table maintained at 125 cm below the surface of the soil. One column was used for the experiment with intermittent application of 2.5 cm of water every five days and the other was used for the experiment with continuous application of 15 cm of water. Applications of water were made with a device constructed from a piece of polyvinyl chloride tube, a piece of lucite, and nine #25 g hypodermic needles. The lucite sheet was glued to the bottom of the polyvinyl chloride tube and the hypodermic needles were glued to the lucite sheet. These needles delivered water on the soil surface as small droplets at a uniform rate maintained by a constant water head in the polyvinyl chloride tube. Five days after application of 2.5 cm of water at 2.5 cm/hr, KCl equivalent to 180 kg K'ha (based on surface area) was mixed in the top 3 cm of the soil and another 2.5 cm of water was applied at the same rate. Soil solution samples were collected after 15 hr. The application of water was repeated three more times at five-day intervals. Soil solution samples were collected each time through each of the ten tensiometer-samplers after 15 hr. All soil solution samplers were flushed with distilled water before soil solution was extracted. The soil solution sampling equipment is shown diagramatically in Figure 2. In the second soil column, where 15 cm of water were applied continuously soil solution samples were collected only at the end of the water application period. All samples were analyzed for K but samples from the experiment with continuous application of water were also analyzed for Ca and Mg.

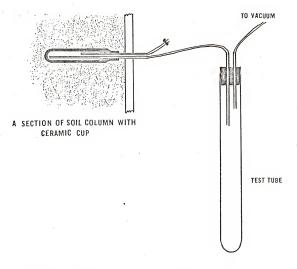


Figure 2. Soil solution sampling apparatus used in the laboratory experiment on K leaching.

Miscible Displacement Studies

To determine K and Cl⁻ dispersion in Lakeland fine sand during saturated or unsaturated flow of a solution containing KCl, experiments were carried out on reconstituted columns of Lakeland fine sand obtained by packing air-dried samples from 0-10, 10-20, and 20-30 cm depths.

For the saturated case, the soil samples were packed into a glass tube 2.5 cm inside diameter and 30 cm long. One pore volume of 0.01 M KCl solution labelled with 0.01 Ci/ml of Cl36 (as NaCl) was applied to the horizontally mounted column and was followed by distilled water. The flow through the soil column was controlled and maintained by a pump. Effluent fractions of 5.5 ml each were collected with a fraction collector starting at the time the KCl solution was introduced into the column. The flow velocities tested were 18, 22, and 69 cm/hr. One pore volume was determined from the actual saturated water content of the soil in the glass tube.

A vertically mounted column was used for the study under unsaturated condition. The apparatus consisted of a soil sample retaining cylinder constructed of perforated stainless steel 10 cm inside diameter and 20 cm long, two stainless steel base plate, and a larger stainless steel cylinder for holding air pressure (Figure 3). Each base plate had two 0-rings fitted for the outer and inner cylinders and a porous plate. The base plates also had centered and off-centered nipples that led to the compartment of the ceramic plate. The centered nipple was used for flushing the space between the ceramic plate and the base plate. The top base plate had two extra nipples which passed into the space between the outer and inner cylinders, one for air pressure and the other to

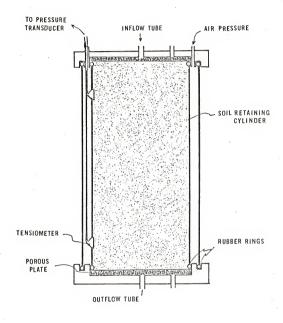


Figure 3. Apparatus used in the miscible displacement study under unsaturated condition.

connect the two tensiometers placed 15 cm apart along one side of the wall of the sample retaining cylinder to a pressure transducer.

After saturating the porous plates under vacuum, the sample retaining cylinder was properly placed on the lower base plate and the soil samples were packed in proper order. After the whole apparatus was assembled the soil was saturated through the bottom plate and allowed 12 hr of equilibration before pressure was introduced for desaturation. During desaturation water was continuously flowing through the soil column at a rate regulated by a pump. The proper rate of flow was determined by trial and error method based on the tensiometer readings of the upper and lower tensiometers. The tensiometers were connected to a pressure transducer and soil water tension was measured potentiometrically after calibration against standard pressure heads. The soil was allowed to attain equilibrium conditions, established as an applied air pressure equivalent 50 cm of water. While water was flowing through the soil at a constant rate, a 0.01 M KCl solution labelled with Cl³⁶ was introduced into the soil column. The effluent was collected in 5.5 ml increments with a fraction collector. All fractions were analyzed for K by the usual procedure and for $C1^{36}$ activity using a Tri-Carb liquid scintillation counter.

Miscellaneous Analyses

The K-Ca exchange characteristic of the soil was determined by quantity-intensity (Q/1) technique proposed by Beckett (1967). Samples were equilibrated with 50 ml of 0.001 $\underline{\text{M}}$ CaCl₂ solutions of varying KCl concentrations ranging from 0 to 0.002 $\underline{\text{M}}$. Equilibration with the solution without K was performed for 0.5, 1.5, 4.5, and 7.0 g samples in 50 ml of

the solution, to determine K release characteristic. For the other solutions (with K) equilibrations were performed using 7.0 g soil samples. The equilibrium solutions were analyzed for Na and K by flame spectroscopy and for Ca and Mg by atomic absorption spectroscopy. The gain or loss of K in the equilibrium solution from that of the original solution was calculated as well as the activity ratio of K to (Ca+Mg) by the use of the Debye-Huckel equation for correcting activities of ions.

Organic matter content of soil samples were determined by the Walkley and Black method (Allison, 1965). Cation exchange capacities were determined from 25 g soil samples by repeated leaching with $\underline{\text{N}}$ NH₄OAc and then replacement of NH₄ on the soil exchange by leaching with $\underline{\text{N}}$ NaCl. The NaCl leachate was analyzed for N by a micro-Kjeldahl distillation procedure, with CEC determined from the amount of NH₄ replaced by Na.

Water retention characteristics of the soil samples were determined in the soil physics laboratory in the Department of Soil Science of the University of Florida, on undisturbed core samples. The core samples were mounted in Tempe cells through which water retention at various applied pressures was determined gravimetrically. A pressure membrane extractor was used for higher applied pressures. Saturated hydraulic conductivity was determined on the same samples and used to calculate hydraulic conductivities at different water contents were calculated following the procedures given by Green and Corey (1971). Calculations were all made with the help of an IBM 360 computer. The plots of water content vs applied pressure were used to determine water contents corresponding to tensiometer readings.

RESULTS AND DISCUSSION

Field Experiments

Field Experiments on Flue-Cured Tobacco

In this manuscript, the term "treatment" réfers to fertilizer material or method of application such as for 4-3.5-10 which was applied either split or once in the 1972 experiment. Therefore, the rates of application are not referred to when treatment effects are discussed. Either the fertilizer name or the corresponding treatment number will be used. Complete outlines of the two field experiments may be found in Tables 1 and 2.

<u>Field experiment-1972</u>. The yield; K, total alkaloid, and total reducing sugar contents of the leaves; price per lilogram weight of the cured leaves; and their analysis of variance for the 1972 field experiment on flue-cured tobacco are given in Tables 3 to 16. In the analysis of variance tables, comparison between treatment 8, where all N was applied at planting, and the second rate of treatment 5, where N was applied split, is denoted by "S5, R2 vs S8". The comparison between the check treatment (no K applied) and all other treatments except 8 is denoted by "Ck vs others".

Table 3 contains the yield data by priming and the analysis of variance for these data are in Table 4. Treatments did not result in significant yield differences at any of the five primings but at every priming, except the first, differences in fertilizer rate were significant. In the overall analysis of variance (Table 5), treatment 8 in which all N was applied at planting was significantly lower in yield than treat-

Table 3. Mean yield of flue-cured tobacco in 1972 experiment.

Treat- ment	Treatment1/	Rate	2/		Primin	g		Total
No.	77 55 21110110	- Na CC	1	2	3	4	5	iotai
1	4-3.5-10 (split)	1 2 3	330 379 324	294 328 333	302 345 392	y/ha 431 445 571	551 473 717	1908 1970 2338
2	4-3.5-10 (once)	1 2 3	315 324 300	284 351 291	339 283 375	389 425 349	540 456 619	1867 1839 19 3 4
3	SCK (36% coating)	1 2 3	342 336 325	294 335 334	358 385 412	388 466 500	411 493 546	1793 2015 2117
4	SCK (18% coating)	1 2 3	329 337 370	308 362 326	315 351 445	424 465 451	488 599 636	1864 2114 2228
5	0 smocote (7-6-17.5)	1 2 3	348 368 331	361 345 250	361 366 444	362 432 611	511 592 569	1943 2103 2305
6	Osmocote N	1 2 3	348 391 384	208 400 342	303 351 417	399 369 417	372 295 713	1630 1806 2273
7	4-3.5-0		331	359	348	484	568	2090
8	Osmocote (7-6-17.5)	2	352	354	262	334	307	1609

 $^{1/\}text{See Table 1 for details.}$

 $^{2^{\}prime} {\rm Rates}$ 1, 2, and 3 correspond to 450, 900, and 1800 kg/ha of 4-3.5-10, respectively.

Table 4_\bullet . Analysis of variance for yields of flue-cured tobacco from 1972 experiment

Source of	J.			Priming			Takal
variation	;	_	2	3	4	5	iorai
				mean square -			
Blocks	2	13653**	14514**	23081**	20621	115662**	280854**
Treatments (S) Rate	2	2829	1648	3581	10793	17631	68832
Linear	-	-	5629	56299≈≈	20994**	199436**	**##1296
Quadratic	-	3439	22195**	1123	393	19716	2867
S x Rate	0	842	3330	1368	8391	18213	28140
S5, R2 vs. S8	-	294	112	12788	11440	97282	289960**
Ck vs. others	-	348	2487	227	5843	5158	27029
Error	38	1688	2220	2825	6387	12579	30045

* Significant at the 5% level

Table 5. Overall analysis of variance for yields, K uptake, and K, total alkaloid (TA), total reducing sugar (RS) contents of leaves in 1972 field experiment.

Source of variation	df	Yield	K uptake	%К	%ТА	%RS
			mean	square		
Blocks	2	56170**	143.4%	5.20%	3.05**	1.48
Treatments (S) Rate	5	13766	30.7**	2.12**	0.37*	0.12
Linear	1	193428**	409.3**	9.83%	2.69%	3.63**
Ouadratic	1	573	0.2	0.21	0.29	0.01
S x Rate	10	5628	8.7	0.37	0.37%	
S5, R2 vs S8 (C1)	1	57992**	44.5**	0.31	0.29	0.31
Ck vs Others (C2)	1.1	5405	96.4**	10.78**	4.24%	0.15
Error (a)	38	6009	8.5	0.26	0.12	0.12
Primings (P)	4	317192**	102.5**	37.29**	53.77**	14.99%
P x Treatments	20	5695	4.4	0.26**	0.10	0.06
P x Rate	8	20653***	7.7*	0.23*	0.11	0.36%
P x S x Rate	40	6629	4.3	0.09	0.20%	0.04
P x C1	4	15981	2.7	0.22	0.01	0.11
P x C2	4	2164*	7.7	0.68	0.00	0.07
Error (b)	160	6319	3.5	0.10	0.13	0.07

^{*} Significant at the 5% level ** Significant at the 1% level

ment 5 where 34 kg N/ha was applied five weeks after planting. This suggests that sidedressing of N is beneficial. The yields from the first six treatments did not differ significantly from that of treatment 7 where no K was applied, indicating that K was not limiting tobacco yields in this experiment. Total yields did not vary significantly among treatments (Table 4) but increased linearly with rate of N, P, and K fertilizers. Regression equations for the effect of rate on yields are given in Table 6. Except for the second priming, for which the yield response was quadratic, the responses were linear with rate. These yield responses with rate could not be attributed to K since there was no treatment x rate interaction effect and N and P were also varied with K rate.

The influence of fertilizer treatments and rates on the K contents of the tobacco leaves from each priming are shown in Table 7. The analyses of variance for individual priming data are shown in Table 8. The application of K fertilizer increased the K contents of the leaves from all primings. Increasing rates of fertilizer increased K contents linearly at all primings but the effect was less pronounced in the later primings as indicated by the decreasing value of the regression coefficients given in Table 9. Table 10 shows which treatments significantly influenced K content as determined by Duncan's multiple range tests (DMRT). Treatments 1, 5, and 6 resulted in higher K contents of the leaves than treatments 2, 3, and 4. This indicates that sulfur-coated K₂SO₄ (treatment 3 and 4) possibly had lower solubility thus restricting K availability during periods of high K absorption by the tobacco plants. Treatment 2, where 4-3.5-10 was applied once at planting, had the lowest K in the leaves possibly as a result of K leaching.

The overall analysis of variance for K uptake, the product of yield

Table 6. Regression equations for yield and K uptake based on rate.

Response	Priming		Ε	quation1/
Yield	2	180.8	+	135.3 _R - 24.5 R ²
Yield	3	295.9	+	29.0 R
Yield	L ₄	374.2	+	27.6 R
Yield	5	405.1	+	54.6 R
Total Yield		1722.6	+	120.2 R
K uptake	1	10.4	+	0.7 R
K uptake	2	4.2	+	4.7 R - 0.7 R ²
K uptake	3	7.5	+	1.5 R
K uptake	4	6.9	+	1.1 R
K uptake	5	5.2	+	1.3 R
Total K uptake		38.1	+	5.5 R

 $[\]frac{1}{R} = \frac{\text{rate of fertilizer applied (kg/ha) equivalent to } 4-3.5-10.}{450}$

Table 7. Mean K contents of leaves in 1972 experiment.

Treat	- 1/	- 2/			Priming		
ment No.	Treatment 1/	Rate ² /	1	2	3	4	5
1	4-3.5-10 (split)	1 2 3	3.5 3.5 4.5	2.8 2.7 3.9	2.8 3.2 4.0	2.0 2.1 2.7	1.4 1.5 1.7
2	4-3.5-10 (once)	1 2 3	3.5 3.5 4.0	2.9 2.6 3.3	2.8 2.7 2.9	2.2 2.3 2.4	1.4 1.3 1.6
3	SCK (36% coating)	1 2 3	2.7 2.8 3.7	2.2 3.1 2.9	2.5 2.8 2.9	1.9 2.0 2.2	1.4 1.4 1.5
4	SCK (18% coating)	1 2 3	2.8 2.9 3.4	2.5 3.1 2.6	2.7 2.8 3.3	1.8 1.9 2.1	1.4 1.4 1.5
5	Osmocote (7-6-17.5)	1 2 3	3.6 3.7 4.1	2.9 3.6 3.7	3.0 3.4 3.3	1.9 1.9 2.3	1.4 1.5 1.5
6	Osmocote N	1 2 3	3.8 3.5 4.1	3.4 3.0 3.6	3.2 3.1 3.5	2.1 2.2 2.3	1.8 1.6 1.9
7	4-3.5-0		1.9	2.1	2.2	1.7	1.1
8	Osmocote (7-6-17.5)	2	3.2	2.9	3.2	2.2	1.6

 $^{1/\}mathrm{See}$ Table 1 for details. $2/\mathrm{Rates}$ 1, 2, and 3 are equivalent to 450, 900, and 1800 kg/ha of 4-3.5-10.

Table 8. Analysis of variance for K contents of the leaves in 1972 experiment.

Source of	df			Priming		
variations	u i	1	2	3	4	5
				mean squ	are	
Blocks	2	1.19%	2.32**	1.28**	1.07**	0.34**
Treatments (S)	5	0.32***	0.83*	0.58%	0.22%	0.20%
Rate						
Linear	1	4.5 h/c/c	2.79**	2.33%%	1.14%	0.37**
Quadratic	1	0.69	0.02	0.00	0.00	0:04
S x Rate	10	0.09	0.42	0.16	0.04	0.02
\$5, R2 vs \$8	1	0.35	0.68	0.08	0.13	0.01
Ck vs others	1	7.46%	2.70%	2.29%	0.59%	0.46%
Error	38	0.17	0.23	0.12	0.05	0.02

^{*} Significant at the 5% level. ** Significant at the 1% level.

Table 9. Regression equations for K contents of the leaves based on rate.

Priming	Equation 1/
1	2.99 + 0.23 R
2	2.61 + 0.18 R
3	2.67 + 0.16 R
4	1.87 + 0.11 R
5	1.35 + 0.06 R

 $[\]frac{1}{R}$ = $\frac{\text{rate of fertilizer applied (kg/ha) equivalent to 4-3.5-10}}{450}$.

Table 10. Results from Duncan's multiple range test for treatment means in 1972 experiment.

Response1/	Priming		Tr	eatmen	t numb	er <u>2</u> /	
% к	1	1	6	5	2	3	
% к	2	5	6	11	2	3	- 1
% к	3	1	6	5	4	2	:
% к	4	2	1	6	5	3	ı
% к	5	<u>6</u>	1	_5	2	4	
K uptake	1	6	1	5	2	4	:
K uptake	3	5	6	1	4	3	:
Total K upt	ake	5	1	6	4	2	-
% TA	2	3	4	2	1	5 -	

 $[\]frac{1}{2}/$ Responses with significant effect of treatments. $\frac{2}{2}/$ Treatment numbers underscored with same line mean that effects are not significantly different.

and K content, was included in Table 5. Application of K significantly increased K uptake at the first and third priming, and in the total of the five primings. Results from DMRT on treatment means (Table 10) showed higher total K uptake from treatments 1, 5, and 6 than treatments 2, 3, and 4 similar to that observed for K content of the leaves. The response to rate of fertilizer application was linear at every priming except at the second where the response was quadratic (Table 6).

The total alkaloid and total reducing sugar contents of the leaves from each of the five primings are given in Table 11. The analyses of variance are given in Table 12. Neither total alkaloid nor reducing sugar content of flue-cured tobacco varied significantly among treatments except in the second priming. The total alkaloid content increased linearly with rate of fertilizer application from first to fourth priming. Variations between treatment 7 and the other treatments were significant suggesting that higher K levels may be involved in keeping the total alkaloid content of the leaves at a low level. The regression equations based on rate of application for the first, second, and fourth primings are given in Table 13.

Total alkaloid content increased with priming. This is consistent with the general observation with tobacco that nicotine (an alkaloid) increased from the bottom to the top of the tobacco plant (Hawks, 1970). On the other hand, there was no consistent trend for reducing sugar content with regards to priming except that it is highest in either fourth or fifth priming.

The overall analysis of variance (Table 5) indicated a significant priming x treatment x rate interaction for total alkaloid contents.

Table 14 shows the mean total alkaloid content from five primings for

Mean total alkaloid (TA) and total reducing sugar (RS) contents of flue-cured tobacco leaves in 1972 experiment. Table 11.

	/lacatecat	2/					- I	Priming				
No.	1	kate-	TA	RS	TA	2 RS	TA	3 RS	ΔT	4 RS	T.	5
-	4-3.5-10 (split)	_	1.43	10.7	1,45	11.4		 %	2 63	0 01	2 28	200
	=	2	1.29	9.2	1.57	8	3.00	9	3.5	17.7	3.6	٠-
	=	3	1.58	4.7	1.62	5.4	2.81	2.0	2.81	17.2	3.32	
2	4-3.5-10 (once)	-	1 22	,	1 22	10 1	1	1	ć	1	1	
ı	=======================================	- c	00.		200	7.7	2.7	٥٠/	7.88		3.78	
	=	7 6		۔ . ن	79.1	- 1	2.48	7.9	2.91	9.91	3.33	
		^	.54	٥.٥	1.73	1.7	3.04	5.9	3.30	14.2	3.67	17.8
~	SCK (36% coating)	-	1.36	7.6	1 43	15 3	2 91	2	37 6	01	07.0	-
	=	2	1.42	0	1.75	000	2 73	9	3 23		2 77	
	=	6	1.69	5.2	1.86	7.4	2.95	9.0	3.72	13.5	3.59	8
4	SCK (18% coating)	-	1.24	12.3	1.30	14.3	2.83	8.9	2.57	20.1	3.59	-
	=	2	1.58	8.6	1.57	9.5	2.61	7.2	3,26	16.4	3.56	18.7
	=	m	1.61	4.9	1.89	8.4	2.52	4.5	3.45	17.2	3.15	-
2	Osmocote (7-6-17.5)	_	1 38	0	1 4.7	11 3	2 111	7	,	2 21	1	•
	= =	2	1.16		3	, ,	2 60	, 4	20.0	2 0	2.13	
	=	٣	1.38	5.0	1.53	8.1	2.50	5.5	2.84	18.0	3.32	19.2
9	Osmocote N	-	0.95	8	1.03	12.8	2 23	~	3 12	18 2	200	_
	Ξ	2	1.16	1.4	7		200	· «	2.5	16.2	200	
	=	~	1.68	6.9	1.48	4.6	2.85	6.7	2.94	18.0	4.15	16.7
7	4-3.5-0		1.86	10.5	1.98	7.4	3.17	9.9	3.61	15.3	4.08	16.9
œ	Osmocote (7-6-17 E)	¢	-			7.	c	c				
•		4	. 03	6.2	50.1	4.5	7.30	0.0	7.65	4.6	2.99	4.

<u>1</u>/See Table 1 for details. $\frac{2}{\text{Rates}}$ 1, 2, and 3 are equivalent to 450, 900, and 1800 kg/ha of 4-3.5-10, respectively.

Table 12. Analysis of variance for total alkaloid and reducing sugar contents of tobacco leaves from 1972 experiment.

Source of	df			Priming		
variations		1	2	3	4	5
			m	ean squar	e	
			Tot	al Alkalo	ids	
Blocks	2	0.65**	0.45**	0.55	0.22%	2.27%
Treatments (S) Rate	5	0.07	0.23**	0.24	0.14	0.09
Linear	1	0.70%	1.10%	0.62**	0.88%	0.00
Quadratic	1	0.01	0.02	0.05	0.41	0.11
S x Rate	10	0.08	0.04	0.37%	0.30	0.39
S5, R2 vs S8	1	0.01	0.10	0.07	0.02	0.16
Ck vs others	1	0.67**	0.72**	0.96**	0.98*	0.92
Error	38	0.05	0.04	0.11	0.20	0.24
			Red	ducing Su	ars	
Blocks	2	0.27	0.72**	0.47**	0.43%	0.17%
Treatments (S) Rate	5	0.13	0.13	0.02	0.06	0.02
Linear	1	2.36***	2.26**	0.43**	0.33	0.00
Quadratic	1	0.66	0.23	0.00	0.26	0.00
S x Rate	10	0.06	0.06	0.02	0.08	0.05
S5, R2 vs S8	1	0.03	0.67*	0.07	0.01	0.01
Ck vs others	1	0.06	0.25	0.04	0.10	0.05
Error	38	0.09	0.09	0.06	0.09	0.04

^{*} Significant at the 5% level. ** Significant at the 1% level.

Table 13. Regression equations for total alkaloid (TA) and reducing sugar (RS) contents of tobacco leaves based on rate.

Response	Priming	Equation 1/
% ТА	1	1.17 + 0.09 R
% TA	2	1.23 + 0.11 R
% TA	. 4	2.80 + 0.10 R
% RS	1	0.69 + 0.44 R - 0.12 R
% RS	2	1.40 - 0.16 R
% RS	3	0.86 - 0.07 R

 $[\]frac{1}{8}$ R = $\frac{\text{rate of fertilizer applied (kg/ha) equivalent to 4-3.5-10}}{450}$.

Table 14. Treatment x rate interaction means for total alkaloid contents of tobacco leaves in 1972 experiment.

Treat-		Ratel/		2/
ment No.	1	2	3	Equation2/
		%		
1	2.34ab <u>3</u> /	2.59a	2.32bc	$1.83 + 0.63R - 0.13R^2$
2	2.22ab	2.35ab	2.63a	2.08 + 0.14 R
3	2.39a	2.58a	2.76a	2.29 + 0.12 R
4	2.24ab	2.53a	2.58ab	2.21 + 0.10 R
5	2.43a	2.23b	2.29c	No significant regression
6	2.05b	2.36ab	2.62a	1.92 + 0.18 R

 $[\]frac{1}{2}$ Rates 1, 2, and 3 correspond to 450, 900, and 1800 kg/ha of 4+3.5-10, respectively.

 $[\]underline{2}/R = \frac{\text{rate of fertilizer applied (kg/ha) equivalent to } 4-3.5-10}{450}$.

^{2/} DMRT on treatment means (comparisons within a rate). Values with same letters are not significantly different.

various treatments and rates of application and the regression equations for each treatment based on rate of application.

The total reducing sugar content decreased linearly with rate of application in the second and third priming while response with rate in the first priming was quadratic. The overall analysis of variance (Table 5) indicates that the relationship between rate of application and reducing sugar content is linear.

The grade of flue-cured tobacco is an index of the quality of the cured leaves. Although the market price for a particular grade may vary from year to year, the relative values of the different grades remain about the same. Hence, for statistical purposes the relative prices for the season would be indicative not only of quality for the season but also of the relative values of the grades for any season.

The data on price per kilogram weight of flue-cured tobacco in the 1972 experiment are given in Table 15. Analyses of variance for each of the five primings and that for the total value are given in Table 16. The price of flue-cured tobacco did not vary among treatments. Increasing rate of fertilizer application increased price linearly in the first and third priming. The total value, determined from the products of price and yield from each priming, increased significantly with rate of application.

The data on K, Ca, and Mg content of the soil samples representing four depths and five samplings are given in Tables 17, 18, and 19, respectively. The analyses of variance of the data are given in Table 20. Significant differences due to treatments were observed for K, Ca, and Mg contents of the samples. The effect of treatments on K was independent of sampling time. The DMRT results showed that the soil samples

Table 15. Prices of flue-cured tobacco in 1972 experiment.

Treat		Rate2/			Primin	g	
No.	Tredement.	1/4 CC	1	2	3	4	5
1	4-3.5-10 (split)	1 2 3	1.84 1.84 1.83	1.88 1.88 1.86	- \$/kg 1.91 1.91 1.88	1.88 1.88 1.91	1.89 1.89 1.89
2	4-3.5-10 (once)	1 2 3	1.84 1.85 1.84	1.87 1.87 1.87	1.91 1.92 1.88	1.88 1.91 1.88	1.91 1.88 1.91
3	SCK (36% coating)	1 2 3	1.85 1.86 1.83	1.89 1.89 1.89	1.91 1.91 1.91	1.91 1.91 1.91	1.88 1.87 1.87
4	SCK (18% coating)	1 2 3	1.85 1.85 1.84	1.89 1.88 1.91	1.89 1.88 1.91	1.81 1.91 1.91	1.89 1.91 1.91
5	Osmocote (7-6-17.5)	1 2 3	1.85 1.85 1.83	1.88 1.86 1.87	1.91 1.91 1.87	1.89 1.88 1.85	1.88 1.88 1.89
6	Osmocote N	1 2 3	1.83 1.85 1.83	1.88 1.89 1.87	1.92 1.92 1.91	1.91 1.91 1.85	1.87 1.87 1.88
7	4-3.5-0		1.83	1.88	1.92	1.88	1.88
8	Osmocote (7-6-17.5)	2	1.84	1.89	1.91	1.88	1.89

 $[\]frac{1}{2}$ (See Table 1 for details. $\frac{2}{2}$ (Rates 1, 2, and 3 are equivalent to 450, 900, 1800 kg/ha 4-3.5-10, respectively.

Table 16. Analysis of variance for price of flue-cured tobacco data in 1972 experiment.

Source of	df			Priming			Total
variations		1	2	3	4	5	value
0.1 1	_			mea	n square -		
Blocks	2	1.67*	2.15	3.01	17.12**	4.72*	214023%
Treatments (S) Rate	5	0.59	2.92	1.31	2.09	3.39	50856
Linear	1	3.17**	0.13	5.41*	2.01	2.57	694533
Quadratic	- 1	1.75*	0.02	0.39	1.43	0.01	2905
S x Rate	10	0.22	0.54	1.42	2.41	0.82	19616
S5, R2 vs S8	1	0.17	2.67	0.17	0.00	0.67	207628**
Ck vs others	1	0.59	0.04	1.54	0.50		
Error	38	0.38	2.62	1.19	2.01	0.14 1.45	18759 21527.

Table 17. Effect of treatments on K contents $\frac{1}{2}$ of soil in 1972 field experiment.

Treat-	Treatment2/		Dept		
ment N	0.	0-15	15-30	30-45	45-60
1	4-3.5-10 (split)	63	69	om	26
2	4-3.5-10 (once)	80	144	72	34
3	SCK (36% coating)	129	138	51	24
4	SCK (18% coating)	115	104	60	25
5	Osmocote (7-6-17.5)	102	71	56	19
6	Granular K ₂ SO ₄	118	164	111	22

 $[\]underline{1}/$ Means of five samplings and three rates.

^{2/} See Table 1 for details.

Table 18. Treatment x depth interaction means $1/\sqrt{100}$ for Ca contents of the soil.

	Treat-		Depth3	/		Test
_	ment No.2/	1	2	3	4	results4/
			ppm	Ca		
	1	204	111	92	60	(1) (2 3 4)
	2	256	252	58	43	(1 2) (3 4)
	3	326	168	7 7	45	(1) (2 3) (3 4)
	4	272	225	69	41	(1 2) (3 4)
	5	124	75	40	26	(1 2 3 4)
	6	169	93	61	34	(1 2) (2 3 4)

 $[\]frac{1}{2}$ /Means of five samplings and three rates. $\frac{2}{3}$ /See Table 1 for details.

² Depths 1, 2, 3, and 4 correspond to depths 0-15, 15-30, 30-45, and 45-60 cm, respectively.

^{4/}Numbers in parentheses represent depths. Parentheses denote non-significant differences among values of Ca content for the depths indicated.

Table 19. Treatment x depth ineraction means 1/ for Mg contents of the soil.

Treat-		Dep	th3/		Test	-
ment No.2/	1	2	3	4	results4/	
		рр	m Mg		1030113	-
1	21	13	7	5	(1 2) (2 3) (3	4
2	19	18	9	6	(1 2) (3 4)	
3	29	32	20	6	(1 2) (3) (4)	
4	32	29	21	7	(1 2) (2 3) (4)	
5	19	13	9	4	(1 2) (2 3) (3	4)
6	30	26	15	4	(1 2) (3) (4)	

 $[\]frac{1}{2}$ /Means of five samplings and three rates. $\frac{2}{2}$ /See Table I for details.

²⁾ See Table 1 101 details. 2) Depths 1, 2, 3, and 4 correspond to depths 0-15, 15-30, 4,30-45, and 45-60 cm, respectively. 4 Numbers in parentheses represent depths. Parentheses

denote non-significant differences among values of Mg content for the depths indicated.

Table 20. Analysis of variance for K, Ca, and Mg contents of the soil in 1972 experiment.

Source of				
variations	df	K	Ca	Mg
			mean square -	
Blocks	2	82302***	81907	249
Samplings (S)				
Linear	1	258	2416	88
Quadratic	1	1538	890	17
Cubic	1	19971	31641	452
Quartic	1	15302	208	58
Treatments (T)	5	22301*	84253*	1583***
SxT	20	13370	29976	97
Error (a)	58	9011	28173	233
Depth (D)	3	145346**	639895***	6889**
SxD	12	9206	19011	229*
TxD	15	7141	27902	185*
SxTXD	60	8206	14660	117
Error (b)	180	6015	16222	103
LITOI (b)	100	0015	16222	103

^{*} Significant at the 5% level. ** Significant at the 1% level.

from plots treated with 4-3.5-10 applied split and resin-coated 7-6-17.5 (treatments 1 and 5) contained significantly less K than plots treated with granular K_2SO_4 and with N supplied from Osmocote (treatment 6). The low soil K contents with treatment 1 could not be explained by K uptake since with treatment 6, the K uptakes were also high. The insignificant effect of treatment x depth interaction made it difficult to explain this result on the basis of relative K location within the soil depth sampled. The date however indicate high K content at 3-45 cm depth with the uncoated granular K_2 SO $_4$. It may be postulated that K from this fertilizer moved down the soil profile more quickly than K from other fertilizers or treatments. This downward movement did not significantly reduce K contents of the leaves over that of the other treatments. Split application of 4-3.5-10 resulted in relatively lower K contents throughout the soil depth considered while the single application caused higher K contents particularly at 15-30 cm depth. The K contents from both sulfurcoated materials were about twice as great as that from 4-3.5-10 applied split at 0-15 and 15-30 cm depths. Resin-coated 7-6-17.5 seemed to retard K movement to the lower depths more than the other treatments.

Since K retention in the soil is influenced by the ionic composition of the soil solution, the distribution of Ca and Mg in the soil may be useful in explaining the distribution of K in the soil. In this experiment, there seemed to be no relation of Ca or Mg distribution with K distribution in the soil. Treatments caused significant variations in Ca and Mg contents in the soil. Treatments 2, 3, and 4 resulted in greater amounts of Ca in the soil at 0-15 and 15-30 cm depths than treatments 1, 5, and 6. On the other hand, Mg was higher in the first three layers where treatments 3, 4, and 6 were applied. In addition,

a sampling x depth interaction effect was observed to be significant.

Rainfall data for the experimental area from March to August 1972 are given in Table 21. Within the period between the time the fertilizers were applied (April 3) and the fifth soil sampling (May 24), there was a total of 15.05 cm of water added to the soil as rain. This amount fell in fourteen days. The highest amount (5.82 cm) fell on April 23, three days before the first soil sampling, and the second highest (3.50 cm) fell on May 8, two days prior to the second soil sampling. These two rainfalls increased the water content of the soil to a depth of 30 cm as indicated by soil water tension at various depths measured with tensiometers. The plots were irrigated on May 4 with about 2.5 cm of water but this amount did not seem to change the water content at the 30 cm depth. In contrast, the rainfall on April 23 and May 8 increased the water content at the 60 cm depth and even at the 90 cm depth after redistribution. An additional 2.5 cm of water applied on May 24 increased water contents at 15, 30, and 45 cm depths (Table 22).

The data on soil K content, rainfall, and soil water tension during the first five to six weeks after fertilizer application suggest that K from soluble fertilizers may have moved down as far as 60 cm as a result of downward movement of water from rain and irrigation. There were indications that K from slowly soluble K fertilizers did not move as much as it did from soluble fertilizers. The data on K content of the soil may indicate that K from slowly soluble fertilizers could have positional disadvantage in view of the time-dependent position of maximum root activity of tobacco plants. Hall $\underline{\text{et al}}$. (1953) studied root activity of tobacco by injecting P^{32} solution into the soil at various depths and distances from the plant and measured P^{32} uptake at

Table 21. Rainfall data in 1972 at the experimental site.

Date	March	April	May	June	July	August
1	0	т	T (m	0.48	3.10
	Ö	ò	ò	Ö	0	0.76
2 3 4	1.24	Ö	ō	Ö	Ö	0.70
4	0	ő	Ö	Ö	0	ō
5	2.16	Ö	ō	Õ	Ö	Ö
5 6 7 8	0	ō	Ö	ō	Ö	Ö
7	Ö	0	0.23	0	Ö	2.26
8	ō	ō	3.50	Ö	Ö	T. 20
9	Т	1.29	0	ō	Ö	0
10	0	0	Ō	Ó	1.27	Ö
11	0	0	0	0	T	0.23
12	0	0	0	0	0	0
13	0	0	0	0	0	Ö
14	0	0	0.30	0	1.44	0
15	0	0	0.99	0	0	0
16	0	0	T	0	0.10	0
17	0.50	0	0	0	0	0 -
18	0	0	0	1.03	0	0
19	0	0	0.25	6.10	0	0
20	0	0	0.74	5.38	0.25	T
21	0	0	0	Т	0	2.54
22	0.15	0	0	0	0	0
23	0	5.82	0	0	0	0
24	0	0.23	0	0	0	0.30
25	0	T	0	0.89	3.08	0.81
26	0	0.	1.85	2.16	0.56	4.88
27 28	0	0	0	3.53	T	5.67
29	1.60	0	1.35	1.03	0	T
30	6.15		1.03	0 04	0	0
31	4.33	1.70	0.10	0.94	0	0 T
otal	16.13	9.04	10.42	21.06	7.18	20.55

Soil water tension $\frac{1}{2}$ at various depths, before and after rainfall or irrigation during the five-week soil sampling period in 1972.

 Dat			Dep	th (cm)		
 - Vai		15	30	45	60	90
				cm		
April		85	60	51	30	0
11	28	95	68	55	36	6
May	1	83	68	53	36	6
11	3 .	159	68	52	42	12
- 11	42/	183	68	52	44	14
11	3 4 <u>2</u> / 4 <u>3</u> /	111	68	51	40	10
11	5	81	68	53	44	14
: 11	5 8	149	84	53	50	20
11	10	71 1	61	35	30	0
11	12	137	70	27	40	10
11	15	509	78	53	46	16
11	17	163	78	39	46	.16
11	19	143	82	53	50	20
11	22	143	86	37	52	22
11	23	671	96	29	54	24
11	24	71	78	27	56	26
ìi	26	57	78	7	54	24
11	30	75	78		52	22
11	31	99	84	5 3	52	22
	٠.	22	0-1	,	22	

 $[\]frac{1}{2}$ No measurements made prior to April 26.

^{2/}Before irrigation, 3/After irrigation.

various times. They observed a shift in the maximum root activity of tobacco from the top 15 cm to the layer between 15 and 25 cm below the surface, as early as four weeks after planting. Later, the roots at depths between 25 and 45 cm contributed appreciably to the total ${\sf P}^{32}$ uptake of the plant while the contribution by roots in the top 15 cm declined. It is possible that although K from a soluble fertilizer such as granular ${\sf K}_2{\sf S0}_4$ moves quite readily, the shift in maximum root activity is such that the K may be absorbed before it leaves the root zone. However, if rainfall or irrigation was sufficient to move K to depths lower than the root zone, then there is greater possibility that K leaching will significantly reduce K uptake by plants. In such case, a K source with delayed solubility or slowly soluble property may be an effective source of K.

<u>Field experiment-1973</u>. To further study the effect of K movement on flue-cured tobacco, a second experiment was initiated in 1973. The six fertilizer treatments studied are listed and described in Table 2.

Preplant applications of fertilizers were made on March 23, 1973 but due to bad weather conditions, the seedlings were not planted until March 28, 1973. Approximately 1.14 cm of rain fell between the time of fertilizer application and transplanting. Almost 30 cm of rain fell during the week following transplanting. The water table in the experimental area was at or near the surface for several days. Fortunately, the plants survived although their initial growth was probably slightly hampered. Since there was no apparent runoff within the experimental area, it was decided to continue the experiment. Normally, additional fertilizers would be applied to replace estimated leaching losses but in view of the objective of this experiment, additional fertilizer applications were not made except those originally scheduled such as the sidedressing of NH4NO3 five weeks after transplanting.

Data were obtained for each priming as in the 1972 experiment. Data for yield, K content of the leaves, total alkaloid and reducing sugar contents of the leaves, and the results of statistical analyses are given in Tables 23 to 34. Analyses of variance were performed on each priming and on overall priming data. The overall analysis of variance (Table 25) indicated that fertilizer treatments interacted with priming; therefore, the individual priming analysis of variance provides a better way of studying treatment effects.

Yields varied significantly among fertilizer treatments in the third and fifth primings. The total yield varied significantly among treatments and increased linearly with rate of application. The regression equation

Table 23. Mean yield of flue-cured tobacco in 1973 experiment.

Treat- ment	Treatment1/	Rate	2/	Р	riming			Total
No.			11	2	3	4	5	
1	4-3.5-10 (split)	1 2 3	348 327 342	295 300 357	509 453 465	/ha 342 373 405	508 604 825	2002 2057 2394
2	Swift (0-6.5-25)-IBDU	1 2 3	395 371 299	349 323 333	414 449 450	461 453 421	258 455 529	1877 2051 2032
3	Swift (0-6.5-25)-NH _L NO ₃	1 2 3	290 328 280	306 346 286	432 427 585	464 503 500	689 812 705	2181 2416 2356
4	SCK (26% coating)	1 2 3	319 311 328	284 309 357	354 407 356	436 480 445	588 773 745	1981 2280 2231
5	Osmocote (7-6-17.5)	1 2 3	303 297 379	277 358 294	470 412 386	451 507 528	661 647 580	2162 2221 2167
6	4-3.5-0	1 2 3	258 262 303	307 291 316	447 380 485	460 444 467	573 629 655	2045 2006 2226

^{1/} See Table 2 for details. 2/ Rates 1, 2, and 3 correspond to 450, 900, and 1800 kg/ha of 4-3.5-10, respectively.

Table 24. Analysis of variance for yields of flue-cured tobacco from 1973 experiment.

Source of	¥			Priming			
vatiation	5		2	3	7		Total
				mean square			
Blocks	2	7294*	6598***	31986	57/185	00076	i
Treatments (S)	2	3777	790	11196*	13719	90536	101261
Rates						03000	101701
Linear	-	2454	2556	3484	2611	05053	7000
Quadratic	-	492	1132	4586	2610	2002	200005
S x Rate	10	2265	2337	6360	1634	20052	25033
Error	34	1904	4423	3748	7741	27473	21376

** Significant at the 5% level. ** Significant at the 1% level.

Table 25. Overall analysis of variance for yields, K uptakes, and K, total alkaloid (TA), and total reducing sugar (RS) contents of leaves in 1973 experiment.

Source of variation	df	Yield	K uptake	%K	%TA	%RS
			mear	squares		
Blocks	2	10456	6.24	0.09	1.19**	13.77
Treatments (S)	5	20253**	33.65**	1.08*	0.61*	93.37**
Rate						,,,,
Linear	1	53773**	49.58*	0.11	0.28	27.16
Quadratic	1	11167	2.62	0.14	0.46	5.70
S x Rate	10	5270	13.48	0.53	0.15	21.00
Error (a)	34	4279	7.60	0.38	0.22	20.21
Primings (P)	4	697784**	40.47**	22.02%	42.31**	759.54**
P x Treatment	20	24939**	14.67*	0.39**	0.22*	14.76%
P x Rate	8	11111	7.48	0.09	0.23	4.32
P x S x Rate	40	6140	7.30	0.23*		7.09
Error (b)	144	11460	7.95	0.14	0.12	5.84

^{*} Significant at the 5% level.

** Significant at the 1% level.

for the response of total yield to rates of fertilizer application was: Total yield (kg/ha) = 1995 + 63R. The total yield response due to rate in the 1973 experiment was lower than in 1972.

Results of DMRT (Table 26) showed that the highest total yield was obtained with treatment 3 although it was not significantly different from that obtained with treatment 4, which contained sulfur-coated $K_2 SO_{14}$, nor with treatment 5 which was resin-coated 7-6-17.5. Yields for treatment 6, where no K was applied, did not vary significantly from yields for treatments 1, 2, 4, and 5. In fact, the yields for treatment 6 were slightly higher than yields for treatment 2. Similar to the 1972 experiment, it was not possible to determine whether differences for rates were due to N, P, or K since N and P rates varied with K.

The data on K content of the leaves are given in Table 27. The K content of the leaves varied significantly among fertilizer treatments in the first, second, and fifth primings but the effect in the first priming interacted with rate of application (Table 28). To better understand the nature of the treatment x rate interaction, the regression equations based on rate are given in Table 29 and the results of DMRT on treatment means for each rate are given in Table 30. Treatment 5 resulted in higher K contents in the first priming even at the lowest rate of fertilizer application (Tables 29 and 30). The K content did not vary significantly among rates for treatments 3, 4, 5, and 6. In treatments 1 and 2, the tobacco-grade fertilizer and Swift product 0-6.5-25 with N supplied as IBDU, respectively, the K contentincreased significantly with rate of application. Regardless of rate, the highest K content in the second priming was obtained from treatment 3, the Swift product 0-6.5-25 with N supplied as NH4NO3. In the fifth priming, treatment 5,

Table 26. Results of Duncan's multiple range test on treatment means for yield response in 1973 experiment.

Response <u>l</u> /	Priming		Tre	atme	nt n	umbe	<u>-2</u> /
Yield	3	3	1	2	6	5	4
Yield	5	3	4	1	5	6	2
Total yield		3	5	4	1	6	2

 $[\]frac{1}{}^{\prime}$ Responses shown are those with significant effect of treatments.

^{2/} Treatment numbers underscored with same line do not vary significantly in their yields.

Table 27. Mean K contents of leaves in 1973 experiment.

Treat- ment	Treatment1/	Rate ² /		F	riming		
No.	Treatment—	Kate-	1	2	3 %	4	5
1	4-3.5-10 (split)	1 2 3	2.8 3.4 4.1	2.8 3.1 3.9	2.9 2.8 2.9	2.3 2.2 2.2	2.1 1.5 1.9
2	Swift (0-6.5-25)-IBDU	1 2 3	3.5 3.4 2.5	3.3 3.1 3.2	2.6 2.4 3.2	2.5 2.2 2.4	2.1 2.2 2.2
3	Swift (0-6.5-25)-NH ₄ N	0 ₃ 1 2 3	3.4 3.8 3.1	4.2 3.9 3.8	2.9 2.8 2.7	2.3 2.6 2.1	2.1 2.3 2.2
4	SCK (26% coating)	1 2 3	3.7 3.2 2.9	3.7 3.6 3.3	3.2 2.9 2.8	2.4 2.5 2.4	1.9 2.1 2.1
5	Osmocote (7-6-17.5)	1 2 3	4.0 4.3 4.3	3.7 3.9 3.7	2.8 2.8 2.8	2.6 2.7 2.3	2.3 2.1 2.2
6	4-3.5-0	1 2 3	3.3 2.7 3.7	3.2 3.0 3.7	2.7 2.3 3.1	2.2 2.5 2.6	2.0 1.8 2.1

 $[\]frac{1}{2}/$ See Table 2 for details. $\frac{2}{R}$ Rates 1, 2, and 3 correspond to 450, 900, and 1800 kg/ha of 4-3.5-10, respectively.

Table 28. Analysis of variance for K contents and total K uptakes in 1973 experiment.

Source of	df			Primin	q		Total K
<u>variation</u>		1	2	3	4	5	uptake
				mea	n square -		
Blocks	2	0.95	0.71	0.11	0.01	0.30%	ö≒ 31.23
Treatments (S)	5	1.29**	0.95	0.09	0.11		≈168.23**
Rate						0.21	100.25
Linear	1	0.00	0.22	0.10	0.04	0.02	247.89%
Quadratic	1	0.01	0.07	0.43	0.05	0.07	13.12
S x Rate	10	0.77%	0.32	0.21	0.09	0.08	67.41
Error	34	0.30	0.26	0.17	0.07	0.05	38.02

^{*} Significant at 5% level. ** Significant at 1% level.

Table 29. Treatment x rate interaction means and regression equations for K contents.

Response1/	Treatment		Rate <u>2</u> /		Regression equation3/
nesponse_	No.	1	2	3	Regression equation
% к (1)	1 2 3 4 5	4.1 2.5 3.1 2.9 4.3 3.7	3.4 3.4 3.8 3.2 4.3 2.7	2.8 3.5 3.4 3.7 4.0 3.2	2.5 + 0.4 R 3.9 - 0.3 R Not significant
K uptake (1) kg/ha	1 2 3 4 5	14.3 7.6 8.6 9.8 16.2	11.2 12.6 12.5 9.9 12.7 7.1	9.9 10.3 9.8 11.5 12.0 8.8	8.4 + 1.5 R 4.8 + 7.1R- 1.6 R ² Not significant """ 10.3 + 1.4 R Not significant
K uptake (3) kg/ha	1 2 3 4 5	13.4 14.5 15.8 10.0 10.7 14.9	13.0 10.6 11.9 11.9 11.8 8.6	14.9 10.7 12.3 11.2 12.9 12.1	Not significant 8.7 + 1.4 R 10.3 + 1.3 R Not significant " " 19.8 + 10.0R + 2.1R

 $[\]frac{1}{2}/$ Numbers in parenthesis refer to priming. Rates 1, 2, and 3 correspond to 450, 900, and 1800 kg/ha of 4-3.5-10, respectively. 3/ R = rate of fertilizer applied (kg/ha) equivalent to 4-3.5-10.

⁴⁵⁰

Table 30. Results of Duncan's multiple range test on treatment means for responses with treatment x rate interaction effects.

Response1/	Rate <u>2</u> /	Treatment number ^{3/}
	1	5 4 2 3 6 1
% K (1)	2	5 3 1 2 4 6
	3	5 1 6 3 4 2
	1	5 4 2 1 3 6
K uptake (1)	2	5 2 3 1 4 6
	3	5 1 6 4 3 2
	1	1 5 3 6 4 2
K uptake (3)	2	1 4 3 5 2 6
	3	<u>3 6 2 1 5 4</u>
	1	3 5 1 4 6 2
Price (5)	2	3 4 6 1 2 5
•	3	1 3 4 2 5 6

 $[\]frac{1}{}$ Numbers in parenthesis refer to priming. $\frac{2}{}$ Rates 1, 2, and 3 correspond to 450, 900, and 1800 kg/ha of 4-3.5-10, respectively.

^{2/} Same as those given in Table 2.

Osmocote (7-6-17.5), resulted in the highest K content. This suggests that after heavy rains which occurred during the early part of this experiment, more K was still available to the plant whereas K from other fertilizers, particularly the tobacco-grade fertilizer, had leached beyond the effective root zone.

Where no K was applied, the K content of the leaves did not appreciably differ from those of other treatments. Possibly, lateral movement of water in the soil within the experimental area carried considerable amount of K towards the no-K plots. This possibility is supported by the fact that a drainage ditch adjacent to the experimental area may have provided a sufficient hydraulic head to cause lateral water movement during the period of heavy rainfall. The K content of the leaves from the no-K treatment in the 1972 experiment were lower than those in 1973.

The data on K uptake may not evaluate the availability of slowly soluble K fertilizers since K uptake is a function of yields which may have been more dependent on N. The K uptake varied significantly among treatments in the first and fifth priming. The fertilizer treatments interacted with rates in the first and third priming (Table 29). Rates of treatment 1 also increased K uptake linearly. The highest rate of treatment 2 significantly decreased K uptake, thus a quadratic relation was obtained. Potassium uptake for treatment 3 also followed the same trend with rate as treatment 2 except that the regression was not significant. For treatments 2 and 3, the effect on K uptake in the third priming differed from those of the other treatments. Uptake increased linearly with rates of treatments 2 and 3 but for the same rate, higher K uptake was obtained from treatment 3 that treatment 2. Undoubtedly, this was due to the higher yields obtained from treatment 3, possibly

because $\mathrm{NH_4NO_3}$ was a better source of N than IBDU. The application of DMRT (Table 31) confirmed the higher K uptakes from treatment 3 in the fifth priming. Total K uptake also varied significantly among treatments. Treatments 3, 4, and 5 resulted in higher total K uptakes than other treatments. Potassium uptakes from treatments 3 and 5 were significantly higher than those from treatments 1, 2, and 6. This means that except where IBDU was applied as the source of N, the total K uptake was significantly increased when slowly soluble K fertilizers were used. Furthermore, total K uptake increased linearly with rate of fertilizer application according to the equation: Total K uptake (kg/ha) = 53.5 + 1.9R.

The data on total alkaloid and total reducing sugar content of the leaves are shown in Table 32. Generally, both chemical characteristics of tobacco are lower in the 1973 experiment than in 1972 particularly in the lower leaves which were removed in the first priming. Percent total reducing sugar varied significantly among treatments in the second, third, fourth, and fifth primings while total alkaloid varied only in the second priming (Table 33). The effect of rate was significant only in the first priming for total reducing sugar content and in the fifth priming for total alkaloid content (Table 33) but the general trends with rate were consistent with those in the 1972 experiment, where the total alkaloid content increased with rate of application and total reducing sugar content decreased with rate of application. Treatments 1 and 2 generally caused higher total reducing sugar content in the leaves (Table 34). Total reducing sugar content is reduced by high N content in the plant. Since N from tobacco-grade fertilizer was soluble, leaching of N from this fertilizer probably favored the higher content of total reducing sugar. The slow solubility of N from IBDU in treatment 2 probably reduced the N

Table 31. Results of Duncan's multiple range test on treatment means for K content and K uptake responses in 1973 experiment.

Response1/ P	riming		Trea	atme	nt ni	umbe	<u>-2</u> /
% к	2	3_	5	4	6	1	2
% K	5	5	2	3	4	6	1
K uptake	5	3	4	_5_	6	_1_	2
Total K up	take	3_	5	4	1	6	_2

 $[\]frac{1}{2}$ Responses shown are those with significant effect of treatments.

^{2/} Treatment numbers are same as those given in Table 2.

Table 32. Mean total alkaloid (TA) and total reducing sugar (RS) contents of flue-cured tobacco in 1973 experiment.

Treat-							Prin	Priming				
ment	Treatment $1/$	Rate2/							7	_		
No.			ΤA	RS	TA	RS	TA	RS	TA	RS	TA	RS
				-	-		%	9				
-	4-3.5-10 (split)	-	0.55	3.8	1.38	9.6	2.33	6.6	2.75	11.9	2.74	13.3
	=	2	0.97	6.1	1.47	9.3	2.47	4.6	2.78	14.1	1.98	17.7
	=	~	0.97	3.6	1.43	7.4	2.22	10.5	2.87	14.3	2.50	16.7
2	Swift (0-6.5-25)-1BDU	_	0.82	5.7	1.18	10.8	1.94	14.9	2,34	13.1	3.01	13.4
	=	2	0.77	4.2	1,27	6.6	1.85	15.3	3,14	=	2.46	3.8
	=	~	69.0	4.0	1.39	8.3	2.00	11.6	2.63	9.11	3.36	10.9
~	Swift(0-6.5-25)-NH1,NO,	-	0.75	2.8	1.32	3.2	2.03	8.1	2.99	8.7	2.50	12.8
		2	0.94	3.1	1.49	3.4	2.35	6.3	2.65	6.6	2.88	11.2
	=	3	0.51	2.0	1.47	3.7	2.05	11.2	3.19	10.7	3.35	8.9
4	SCK (26% coating)	_	0.87	4.4	1.58	4.1	2.62	6.9	2.83	11.3	2.86	13.7
	=	2	0.91	4.4	1.65	9.6	2.24	9.1	2,82	10.5	2.38	15.0
	=	3	1.01	3.5	1.67	6.7	2.06	10.1	2.92	9.7	3.37	12.8
2	Osmocote (7-6-17.5)	_	0.88	3.8	1.72	4.4	2.78	6.5	3.30	8.2	3.07	12.8
	=	2	0.82	3.4	1.53	3.7	2.19	10.0	3.19	6.6	2.30	15.4
	=	3	1.01	0.4	1.80	6.2	2.59	7.6	3.25	10.3	2.75	15.2
9	4-3.5-0	_	46.0	0.9	1.67	9.7	2.34	4.6	2.72	13.0	2.83	12.9
	=	2	0.79		1.68	5.9	2.21	12.4	2.86	7.7	3.09	12.2
	=	3	0.79	9.1	1.70	2.2	2.39	6.2	3.25	4.9	2.65	12.1

. We Table 2 for details. $2.\mbox{Kares}$ i, 2, and 3 correspond to 450, 900, and 1800 kg/ha of 4-3.5-10, respectively.

Table 33. Analysis of variance for total alkaloid and total reducing sugar contents of tobacco leaves from 1973 experiment.

Source of	df			Primin	9	
variation	u i	1	2	3	Į.	5
			п	ean squa	re	
			Tot	al Alkal	oids	
Blocks	2	0.51**	0.16	0.51	0.04	0.76
Treatments (S) Rate	5	0.05	0.22*	0.36	0.49	0.37
Linear	1	0.00	0.11	0.11	0.16	0.53
Quadratic	1	0.03	0.01	0.07	0.01	1.59%%
S x Rate	10	0.07	0.02	0.12	0.22	0.37
Error	34	0.07	0.08	0.17	0.21	0.20
			Red	ucing Su	gar	
Blocks	2	4.20	19.89	33.87	14.47	11.90
Treatments (S) Rate	5	5.43	52.06**	41.61*	26.56*	26.73**
Linear	1	15.31*	12.60	0.07	6.26	3.83
Quadratic	1	0.43	0.61	11.92	0.45	15.95
S x Rate	10	4.29	10.84	14.95	12.12	7.16
Error	34	2.99	9.58	13.04	9.16	6.06

^{*} Significant at the 5% level. ** Significant at the 1% level.

Table 34. Results of Duncan's multiple range test on treatment means for total alkaloid and reducing sugar contents of the leaves.

Response1/	Priming		Trea	etme	nt ni	ımbe	r <u>2</u> /
TA	2	5	1	4	6	3	2
RS	2	2	1	6	4	5	3
RS	3	2	1	6	4	3	5
RS	4	1	2	3	4	5	6
RS	5	1	5	4	2	6	3

 $\underline{1'}$ Responses shown are those with significant effect of treatments. $\underline{2'}$ Treatment numbers are same as those given in Table 2.

available to the plants to a level that also favored a higher content of reducing sugars in the leaves than treatment 3 where NH_4NO_3 was applied. Total alkaloid content was generally higher where Osmocote (7-6-17.5) was applied and lower where the Swift product (0-6.5-25) was applied.

Table 35 gives the price per kilogram of cured leaves from five primings. The analysis of variance is shown in Table 36. Generally, the prices were higher than those in the 1972 experiment. Where no K was applied (treatment 6), tobacco leaves had lower market value than where K was applied. The market values of the cured tobacco leaves were relatively high in 1973 and the differences among grades of the cured leaves were not reflected on the prices. Although there were significant differences in prices for the first, fourth, and fifth primings, the differences do not indicate any advantage of the slowly soluble K fertilizers over the tobacco-grade fertilizer (Table 37).

Soil samples were collected five times at weekly intervals starting one week after planting at depths of 0-15, 15-30, 30-45, and 45-60 cm from plots treated with 4-3.5-10, Swift (0-6.5-25) with N applied as NH4NO3, SCK (26% coating), Osmocote (7-6-17.5), and 4-3.5-0. The soil was generally wet during the sampling period which may have increased contamination of the samples since the four depth samples were obtained from the same hole using one sampling tube.

The K contents of the soil samples are given in Table 38. In general, the K contents of the soil were lower than those in the 1972 experiment. There were significant differences in K contents of the soil among the five treatments studied (Table 39). The plots treated with Osmocote (7-6-17.5) had higher K content than those with other treatments (Table 40). Surprisingly, the plots with 4-3.5-0 treatment had relatively high soil K

Table 35. Prices corresponding to the grades of flue-cured tobacco in 1973 experiment.

Treat	- Treatment1/	Rate2/			Priming	9	
ment No.	ireatment_/	Kate=/	1	2	3	4	5
1	4-3.5-10 (split)	1 2 3	1.85 1.91 1.87	1.95 1.97 1.95	\$/kg 1.98 1.98 1.98	1.96 1.99 2.00	1.99 1.98 2.00
2	Swift (0-6.5-25)-IBDU	1 2 3	1.88 1.89 1.88	1.95 1.95 1.96	1.98 2.00 1.98	1.99 1.99 1.99	1.98 1.98 1.99
3	Swift (0-6.5-25)-NH ₄ NO ₃	1 2 3	1.81 1.81 1.77	1.96 1.95 1.95	2.00 1.99 2.00	1.99 2.00 1.98	1.99 2.00 2.00
4	SCK (26% coating)	1 2 3	1.87 1.87 1.89	1.95 1.96 1.95	1.99 1.99 1.99	1.99 2.00 1.99	1.99 2.00 2.00
5	Osmocote (7-6-17.5)	1 2 3	1.87 1.85 1.87	1.95 1.96 1.96	1.98 1.96 2.00	1.99 1.97 1.97	1.99 1.96 1.98
6	4-3.5-0	1 2 3	1.93 1.81 1.69	1.95 1.96 1.94	1.99 1.99 1.95	1.98 1.96 1.96	1.99 1.99 1.96

 $[\]frac{1}{2}/\mathrm{See}$ Table 2 for details. $\frac{2}{\mathrm{Rates}}$ 1, 2, and 3 correspond to 450, 900, and 1800 kg/ha of 4-3.5-10, respectively.

Table 36. Analysis of variance for price of flue-cured tobacco data in 1973 experiment.

Source of	df			Priming			Total
variation	αr	1	2	3	4	5	value
				mear	square -		
Blocks	2	38.88*	0.13	4.79*	0.72	1.05	5114
Treatments (S)	5	27.86**	0.15	0.86	2.71*	1.37**	81873**
Rate							
Linear	1	33.33*	0.01	0.02	0.19	0.19	201980**
Quadratic	1	0.08	1.59	0.01	0.25	0.25	47850
S x Rate	10	13.90	0.24	1.06	1.13	1.02%	22270
Error	34	7.67	0.38	0.93	1.07	0.33	17062

^{*} Significant at the 5% level. ** Significant at the 1% level.

Table 37. Results of Duncan's multiple range test on treatment means for price of flue-cured tobacco and total value in 1973 experiment.

Response1/	Priming		Tre	eatmo	ent i	numbe	e <u>r2</u> /
Price	1	2	1	4	_5	6	3
Price	4	2	4	3_	_1	5	6
Total valu	e	3_	5	4	1	6	2

 $\frac{1}{2}\!/\!\text{Responses}$ with significant effect of treatments. Treatment numbers are same as those given in Table 2.

Table 38. Effect of treatments on K contents $\frac{1}{}$ of soil in 1973 field experiment.

Treat	-	2/				Sampli		
ment No.	Treatm	nents ² /	Depth	1	2	3	4	5
1	4-3.5-10	(split)	cm 0-15 15-30 30-45 45-60	41 45 32 22	41 32 21 17	ppm K 33 25 17	28 29 30 13	34 37 36 21
3	Swift (0-	-6.5-25)	0-15 15-30 30-45 45-60	53 52 25 24	72 70 20 17	33 35 17 13	20 35 32 12	25 52 81 20
4	SCK (26%	coating)	0-15 15-30 30-45 45-60	49 41 30 33	66 54 36 39	64 51 38 21	26 74 39 23	23 93 42 24
5	Osmocote	(7-6-17.5)	0-15 15-30 30-45 45-60	63 72 45 36	61 122 40 20	58 75 32 24	45 74 34 22	67 74 35 20
6	4-3.5-0		0-15 15-30 30-45 45-60	57 60 39 24	55 42 47 21	38 35 16 12	22 26 21 15	24 31 23 12

 $[\]frac{1}{2}/$ Means of three replications. $\overline{2}$ See Table 2 for details. Treatment numbers are the same as those in Table 2.

Table 39. Analysis of variance on K contents of the soil.

Source of	df			Samplin	q	
variation		1	_2	3	4	5
			me	an squa	re	
Blocks	2	1217*	2463	303	321*	738
Treatments (s) 4	673*	1947	1721*	2561*	1482
Error (a)	8	146	2929	250	506	662
Depth (D)	3	2623%%	8835**	3184%	5133**	3883***
SxD	12	94	1024	168	690%	1043
Error (b)	30	154	1092	138	310	579

^{*} Significant at the 5% level. ** Sifnificant at the 1% level.

Table 40. Results of Duncan's multiple range test for sampling x treatment and sampling x depth interaction effects on K content of the soil.

Sampling1/	Test results ² /
	Treatments
1 .	5 6 3 4 1
3	5 4 6 3 1
	Depths
1	2 1 3 4
2	2 1 3 4
3	1 2 3 4
5	2 3 1 4

Y Samplings with observed significant effect of treatments or depths on K content of the soil. Treatment numbers are the same as those in Table 2. Depths 1, 2, 3, and 4 correspond to depths 0-15, 15-30, 30-45, and 45-60 cm, respectively.

content. Significant differences among depths were found for all five samplings but only the differences in the fourth sampling depended on treatments. Table 41 shows the nature of the treatment x depth interaction in the fourth sampling. The distribution of K in the soil treated with 4-3.5-10 was as homogeneous as the distribution of K in the no-K treated plots. With the other treatments, K contents in the soil were generally highest at about 15-45 cm depth.

One week after transplanting, there were high K contents in the soil where Osmocote (7-6-17.5) was applied. Within the period from fertilizer application to the first soil sampling, about 30.7 cm of rain fell (Table 42), 97% of this amount was received within one week period and about 50% was received in one day (April 4). No soil water tension data were gathered from this experiment since the area was wet most of the time however tensiometers were installed. The high rainfall that occurred probably moved most of the K from the surface layer to lower depths. The slow release property of Osmocote (7-6-17.5) probably prevented immediate loss of K from the surface. The relatively high K content found in the soil of plots treated with 4-3.5-0 presumably was caused by mixing of the soil solution with the nearby plots either internally or by surface runoff or both. It is interesting to note that Osmocote (7-6-17.5) still maintained high K contents in the 0-60 cm layer after those heavy rainfalls. The movement of K to the lower depths with passage of time was indicated by the results of DMRT (Table 40).

Field Experiment on Potassium Leaching

The experiment with intermittent application of water was initiated on July 3, 1973. Immediately after applying the fertilizer, the water

Table 41. Results of Duncan's multiple range test for treatment x depth interaction effects on K content of the soil in fourth sampling.

Treatment		Depth	2/		Test results
⊹ No. ¹ /	1	2	3	4	rest resurts
		ppm	К		
1	28 a <u>3</u> /	29 c	30 ab	13 a	(3 2 1 4)
3	43 a	111 a	55 a	19 a	(2) (3 1) (1 4)
. 4	26 a	74 Ь	39 ab	23 a	(2) (3 1 4)
5	45 a	74 Ь	34 ab	22 a	(2 1) (1 3 4)
6	22 a	26 c	21 b	15 a	(2 1 3 4)

^{1/}Treatment numbers are the same as those in Table 2. 2/Depths 1, 2, 3, and 4 correspond to depths 0-15, 15-30, 30-45, and 45-60 cm, respectively. 3/DMRT on treatment means. Values of K content within a depth

followed by same letter are not significantly different.

Table 42. Rainfall data in 1973 at the experimental site.

Date	March	April	May	June	July	August
,		9 00	0 Cr			5.85
1	0	8.90 0	0	0	0	0.25
2	0	0.63	0	0	0	1.07
2 3 4	0	16.92	0	0	0	0.25
-	0	0	0	0	0	0.69
5 6 7 8	0	0	0	0.79	0	T T
7	0	0.66	0	0.79	0	Ť
0	0.38	3.75	0	T	T	2.03
9	5.18	0	1.22	Ť	1.65	0
10	0.66	0	0	2.80	0	0
11	0.00	0	0	0	0	0
12	0	0	0	Ŧ	2.03	2.13
13	0	0	0	0	0	0
14	0	0	0.63	2.18	0	0
15	0	0	1.44	0	0	3.07
16	0	0	0	0.28	0	0.48
17	0.35	0	0	0.20	0.84	2.89
18	0.55	0	Ö	1.17	2.26	0
19	0	0	Ö	0	2.62	1.04
20	0	0	Ö	1.80	T.02	0
21	0.38	0	Ö	0.56	Ť	•
22	0.50	0	Ö	2.80	o o	
23	Ö	Ö	Ö	T	ő	
24	Ö	Ö	ő	o o	ō	
25	0.63	Ö	1.90	Ö	Ö	
26	0.51	3.30	0.56		Ť	
27	0	1.83	3.30		0.05	
28	Ö	0	0	0	1.70	
29	2.74	Ō	Ö	0.38	1.75	
30	0.25	ō	3.30		1.65	
31	0.14		0.05		1.14	
Total	11.22	35.99	12.40	15.62	15.69	

sprinkler system was operated for 2 hr which delivered a total of 5 cm of water over the excerimental area. The initial soil water content and K concentration of the soil solution at different depths are given in Table 43. The water contents were determined from tensiometer readings and the soil water retention curve. The water content increased with depth to near saturation. One hour after the first application of 5 cm of water, water content at 15 cm depth increased from 15% to 36% while no change was indicated for depths lower than 15 cm. The K concentration of the soil solution did not change at any depth. After 7.5 hr, the K content increased from 5 to 9 ppm at 15 cm depth and the water content increased at 60 cm depth and above. The increase in water content in the soil at 60 cm was either due to the original water in the soil pushed by the applied water or the applied water reached 60 cm depth. Redistribution of water to lower depth was accompanied by a decrease in water content near the surface and increase in K concentration of the soil solution to 150 ppm after 22 hr. There appeared to be no K movement between 22 and 74 hr after application of water. The soil water tension data indicated that probably water moved upward in response to hydraulic gradient which developed due to evaporation of water from the soil surface. The upward movement of water may have caused dilution or displacement of the soil solution at the 15 cm depth, thus decreasing the concentration of K in the soil solution at 15 cm.

Five days after the first application of 5 cm of water, a second application of 2.5 cm of water was made. A rainfall of 1.65 cm occurred 24 hr after the second application. These two additions resulted in an increase in soil solution K concentration at 15 cm depth from 77 to 450 ppm (Table $\frac{1}{4}$). No other increases were observed at any depth. A rain-

Table 43. Soil water tension (h), water content (0), and K content at different depths before and after the first application of water.

Depth	h	θ	K	h	θ	K	h	θ	K
cm		%	ppm	cm	%	ppm	cm	%	ppm
	В	efore		Af	ter l	hr	Aft	er 7.	5 hr
15 30 45 60 75 90	87 64 49 35 19 6	15 20 24 33 35 40	5 1 1 0 1	29 64 49 35 19	36 20 24 33 35 40	5 1 1 0 1	56 61 46 34 19 6	23 21 25 34 35 40	9 1 1 0 1
15 30 45	74 64 47	17 20 25	hr 150 1	90 68 52	er 74 15 18 22	hr 150 0	Aft 99 70 55	er 122 14 18 21	2 hr 77 0
60 75 90	34 17 5	34 35 40	0 1 1	37 22 6	32 35 40	0	40 25 11	31 25 40	0

 $[\]frac{1}{\text{Water applied was 5 cm at 2.5 cm/hr.}}$

Table 44. Soil water tension (h), water content (0), and K content at different depths before 1/ and after the second application of water.

_	Dept	hB	efor	'e	Af	ter 2	hr	Afte	er 28	hr	Afte	er 12	0 hr
_	- op c	h	θ	K	h	θ.	K	h	θ	K	h	9	K
		cm	%	ppm	cm	%	ppm	cm	%	ppm	cm	%	ppm
	15	99	14	77	69	19	ns <u>2</u> /	81	16	450	81	16	525
	30	70	18	0	70	18	11	66	19	1	65	19	0
	45	55	21	1	55	21	- 0	52	23	1	50	24	1
	60	40	31	0	40	31	11	38	31	0	36	32	0
	75	25	35	1	25	35	- 11	24	35	1	23	35	1
	90	11	40	1	11	40	11	10	40	1	7	40	1

1/Water applied was 2.5 cm at 2.5 cm/hr and rainfall of 2/1.65 and 2.03 after 24 and 96 hr, respectively. No soil solution sample collected.

fall of 2.03 cm again occurred after 96 hr which increased soil solution K concentration at 15 cm depth from 450 to 525 ppm after 120 hr.

A total of 11.18 cm of water was added to the soil during the experiment. Because of the insensitivity of tensiometer measurements for determining water contents at low soil water tensions, it was difficult to ascertain exactly, based on water content data, how deep the added water moved. Undoubtedly, the 5 cm of water added the first time moved below 15 cm since the water content at this depth increased to 36%. However, the intermittent application of water with a total of 11.18 cm did not appear to have moved fertilizer K down to the 30 cm depth. Thus K was lagging behind the front of water moving downward. The data on miscible displacement which will be discussed later will provide detailed information about the relationship between movement of water and movement of K in the soil. If the water applied in this field experiment had been tagged, for example with tritium, the tracer wouls precisely indicate the extent of water movement at different times and show how far K was lagging behind.

Table 45 shows the data on soil water tension, water content, and K, Ca, and Mg concentration of the soil solution at different depths before and after continuous application of 15 cm of water at 2.5 cm/hr. In this experiment, the soluble and slowly soluble forms of 7-6-17.5 were compared. The initial water content at various depths particularly near the surface were lower than those in the first experiment. Hence, even if the same amount of water was applied, the effect would probably be different from the first experiment. No soil solution samples were obtained initially because tensiometer readings indicated that the soil was relatively dry and it would have been difficult to collect the

Table 45. Soil water tensions (h), water contents (θ), and movement of K, Ca, and Mg from soluble (S) and Slowly soluble (SS) fertilizers (7-6-17.5) following application of 15 cm of water at 2.5 cm/hr.

Depth	h	θ		K	Ca	1	1	Mg
			S	SS	S	SS	S	SS
cm	cm	%				ppm		
Before a	pplic	ation	/					
15	126	12						
30	100	14						
45	72	16						
60	68	18						
75	53	29						
90	38	35						
At the e	nd of	6 hr	water a	nnlicati	0.0			
	01	U 111		ppricati	OII			
15	38	33	250	135	147	28	41	8
15 30	38 36	33 31	250 52	135 75	147 49	28	41	8
	38 36 34	31	250 52 40	75	49	92	15	24
30	36 34	31 32	52 40		49 32	92 42	15 7	24 12
30 45 60	36 34 50	31 32 27	52	75 32 1	49 32 16	92 42 25	15 7 4	24 12 10
30 45	36 34	31 32	52 40 12	75 32	49 32	92 42	15 7	24 12
30 45 60 75 90	36 34 50 45 30	31 32 27 32 38	52 40 12 1	75 32 1 1	49 32 16 20	92 42 25 17	15 7 4 8	24 12 10 6
30 45 60 75	36 34 50 45 30	31 32 27 32 38	52 40 12 1	75 32 1 1	49 32 16 20	92 42 25 17	15 7 4 8	24 12 10 6
30 45 60 75 90	36 34 50 45 30	31 32 27 32 38	52 40 12 1 1	75 32 1 1 1	49 32 16 20 17	92 42 25 17 29	15 7 4 8 4	24 12 10 6 7
30 45 60 75 90 90 hr aft 15 30	36 34 50 45 30 ter wa	31 32 27 32 38 ater a	52 40 12 1	75 32 1 1 1 1 ion	49 32 16 20 17	92 42 25 17 29	15 7 4 8 4	24 12 10 6 7
30 45 60 75 90 90 hr aft	36 34 50 45 30 ter wa	31 32 27 32 38 ater a	52 40 12 1 1 1 pplicat	75 32 1 1 1 1 ion	49 32 16 20 17	92 42 25 17 29 41 108	15 7 4 8 4 25 34	24 12 10 6 7
30 45 60 75 90 90 hr aft 15 30	36 34 50 45 30 ter wa	31 32 27 32 38 ater a	52 40 12 1 1 1 pplicat 180 191	75 32 1 1 1 1 ion 167 122 90	49 32 16 20 17 83 96 321	92 42 25 17 29 41 108 93	15 7 4 8 4 25 34 87	24 12 10 6 7
30 45 60 75 90 90 hr aft 15 30 45	36 34 50 45 30 ter wa 115 100 88	31 32 27 32 38 ater a 12 14 13	52 40 12 1 1 1 pplicat 180 191 185	75 32 1 1 1 1 ion	49 32 16 20 17	92 42 25 17 29 41 108	15 7 4 8 4 25 34	24 12 10 6 7

 $\underline{1}^{\prime}\mathrm{No}$ soil solution samples collected before water application.

necessary volume of solution for chemical analysis. However, initial K distribution would not be expected to vary much from the first experiment since this experimental area was adjacent to the first experimental area.

The results showed movement of K from soluble fertilizer to depth of 60 cm and to depth of 45 cm with the slowly soluble fertilizer, after 15 cm of water was applied. Further movement was observed after 90 hr resulting from redistribution of water to lower depths. Potassium from both fertilizers moved to as far as 75 cm depth but much greater amounts of the soluble fertilizer were observed to move than for the slowly soluble fertilizer. The data also indicate movement of Ca and Mg downward in the soil but to a lesser extent where the slowly soluble fertilizer was used.

Values of hydraulic head (sum of soil water tension and depth) for each depth indicated that initially, water was moving upwards. At the end of application of water (6 hr), the values of hydraulic head indicated that water was moving downward and continued to move until about 90 hr when the hydraulic head was almost the same for all depths. If this was so, then there was probably no further downward movement of K after 90 hr except perhaps through diffusion. Evaporation of water from the soil surface may result in an upward movement of water and affect the distribution of K in the soil.

Laboratory Experiments

Some chemical and physical properties of Lakeland fine sand used in these experiments are given in Tables 46 to 47. Organic matter content and CEC of the soil decreased with depth (Table 46). Organic matter content could not account for all the CEC of the soil. The inorganic fractions of this soil; 3% clay and 4% silt according to Hammond, Lundy, and Saxena (1967); contributed the balance of the total CEC. The amount of K, Ca, and Mg extractable with \underline{N} $\mathrm{NH}_4\mathrm{OAc}$ at both pH levels were highest in the 0-10 cm depth while Al extractable at pH 4.8 was lowest in the 0-10 cm depth (Table 47).

The saturated water content on the average is 38% by volume. The 15-atm water retention decreased with depth from about 3% to 1.5%. The decrease in water content was probably associated with the organic matter content of the soil. Water retention at tensions lower than 15 atm (15000 cm of water) are given in Table 48. Hydraulic conductivity under water-saturated condition varied inconsistently with depth (Table 46). Values of hydraulic conductivity at higher tensions are given in Table 49.

The quantity-intensity (Q/I) relation measured by the amounts of K released or adsorbed by the soil (ΔK) to or from solutions containing K from 0 to 0.002 \underline{M} in 0.001 \underline{M} Ca are given in Table 50. The activity ratio (AR), $a_{\overline{K}}/(a_{Ca+Mg})^{\frac{1}{2}}$, of the equilibrium solution is given below each value of ΔK . A plot of ΔK vs AR is the usual way of showing Q/I relation for a soil. However, values of ΔK corresponding to the different values of AR for Lakeland fine sand were very small which made it

Table 46. Some physical and chemical properties of Lakeland fine sand.

Depth	Organic matter	Cation exchange capacity	Bulk density	Saturated water content	15-atm water retention	Hydraulic conduc- tivity
cm	%	me/100g	g/cc	%	%	cm/hr
0-10 10-20 20-30- 30-40 40-50 50-60 60-70 70-80 80-90 90+	1.91 1.63 1.32 0.79 0.64 0.55 0.47 0.30 0.26	7.7 6.3 5.9 4.3 3.5 2.5 2.2 1.9 1.8	1.44 1.48 1.48 1.48 1.45 1.45 1.57 1.56 1.54	41.5 38.0 34.8 35.2 40.0 40.8 38.5 35.5 40.0 37.2	2.99 2.68 2.75 2.42 1.99 1.96 1.85 1.72 1.67	19.5 15.8 10.0 18.4 29.0 28.0 20.3 11.2 27.6 27.6

Table 47. Extractable K, Ca, Mg, and Al with 1 \underline{N} NH40Ac(pH 4.8 and pH 7)

cm ppm 0-10 23.9 26.0 100.6 106.5 13.7 12.7 15.0 1.0 10-20 8.2 7.2 49.8 51.4 4.3 3.7 25.2 1.5 20-30 5.3 5.7 33.0 27.8 3.3 2.4 33.5 1.5 30-40 4.6 2.9 25.2 16.1 4.4 2.2 31.2 1.0 40-50 4.6 5.7 21.0 16.1 2.9 2.2 34.2 1.0 50-60 3.9 4.6 23.6 21.6 4.1 2.0 31.2 1.2 60-70 4.3 3.9 16.1 11.9 2.7 1.8 30.5 1.2 70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0			K		a	Mg		Al	
0-10 23.9 26.0 100.6 106.5 13.7 12.7 15.0 1.0 10-20 8.2 7.2 49.8 51.4 4.3 3.7 25.2 1.5 20-30 5.3 5.7 33.0 27.8 3.3 2.4 33.5 1.5 30-40 4.6 2.9 25.2 16.1 4.4 2.2 31.2 1.0 40-50 4.6 5.7 21.0 16.1 2.9 2.2 34.2 1.0 50-60 3.9 4.6 23.6 21.6 4.1 2.0 31.2 1.2 60-70 4.3 3.9 16.1 11.9 2.7 1.8 30.5 1.2 70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	Depth	pH 4.8	pH 7	pH 4.8	pH 7	pH 4.8	рН 7	pH 4.8	рН 7
10-20 8.2 7.2 49.8 51.4 4.3 3.7 25.2 1.5 20-30 5.3 5.7 33.0 27.8 3.3 2.4 33.5 1.5 30-40 4.6 2.9 25.2 16.1 4.4 2.2 31.2 1.0 40-50 4.6 5.7 21.0 16.1 2.9 2.2 34.2 1.0 50-60 3.9 4.6 23.6 21.6 4.1 2.0 31.2 1.2 60-70 4.3 3.9 16.1 11.9 2.7 1.8 30.5 1.2 70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	Cm				рр	m			
20-30 5.3 5.7 33.0 27.8 3.3 2.4 33.5 1.5 30-40 4.6 2.9 25.2 16.1 4.4 2.2 31.2 1.0 40-50 4.6 5.7 21.0 16.1 2.9 2.2 34.2 1.0 50-60 3.9 4.6 23.6 21.6 4.1 2.0 31.2 1.2 60-70 4.3 3.9 16.1 11.9 2.7 1.8 30.5 1.2 70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	0-10	23.9	26.0	100.6	106.5	13.7	12.7	15.0	1.0
30-40 4.6 2.9 25.2 16.1 4.4 2.2 31.2 1.0 40-50 4.6 5.7 21.0 16.1 2.9 2.2 34.2 1.0 50-60 3.9 4.6 23.6 21.6 4.1 2.0 31.2 1.2 60-70 4.3 3.9 16.1 11.9 2.7 1.8 30.5 1.2 70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	10-20	8.2	7.2	49.8	51.4	4.3	3.7	25.2	1.5
40-50 4.6 5.7 21.0 16.1 2.9 2.2 34.2 1.0 50-60 3.9 4.6 23.6 21.6 4.1 2.0 31.2 1.2 60-70 4.3 3.9 16.1 11.9 2.7 1.8 30.5 1.2 70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	20-30	5.3	5.7	33.0	27.8	3.3	2.4	33.5	1.5
50-60 3.9 4.6 23.6 21.6 4.1 2.0 31.2 1.2 60-70 4.3 3.9 16.1 11.9 2.7 1.8 30.5 1.2 70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	30-40	4.6	2.9	25.2	16.1	4.4	2.2	31.2	1.0
60-70 4.3 3.9 16.1 11.9 2.7 1.8 30.5 1.2 70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	40-50	4.6	5.7	21.0	16.1	2.9	2.2	34.2	1.0
70-80 3.9 3.6 20.6 12.9 3.8 2.0 30.0 1.2 80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	50-60	3.9	4.6	23.6	21.6	4.1	2.0	31.2	1.2
80-90 3.6 2.8 24.2 10.3 3.4 1.6 28.5 1.0	60-70	4.3	3.9	16.1	11.9	2.7	1.8	30.5	1.2
1.0 20.5 1.0	70-80	3.9	3.6	20.6	12.9	3.8	2.0	30.0	1.2
	80-90	3.6	2.8	24.2	10.3	3.4	1.6	28.5	1.0
90+ 3.6 3.6 29.4 30.4 4.2 2.0 25.0 1.0	90+	3.6	3.6	29.4	30.4	4.2	2.0	25.0	1.0

Table 48. Moisture retention characteristics of Lakeland fine sand. $\hfill \hfill$

:		De	esorp	tion	pressu	re (c	em)			
: 3.5	: 20	: 30	45	: 60	· 80 :	100	: 150 :	200	: 345	15000
				% by	volume					
41	40	40	35	27	19	16	13	13	11	4
39	37	36	30	21	16	14	11		9	Ĺ.
35	35	28	21	16	14	12	11		_	
37	36	34	25	19	15				7	4
40	39	35	26	20	15				á	3
41	40	37	33	28					7	3
39	39	39	35	26	16				7	3
36	36	36	32	26	16				á	3
40	39	37	33	21	12		7		5	3
38	37	37	32	21	12	10	7	6	5	2
	3.5 41 39 35 37 40 41 39 36 40	41 40 39 37 35 35 37 36 40 39 41 40 39 39 36 36 40 39	3.5 : 20 : 30 41	3.5 : 20 : 30 : 45 41 40 40 35 39 37 36 30 35 35 28 21 37 36 34 25 40 39 35 28 41 40 37 33 39 39 39 35 36 36 32 40 39 37 33	3.5 : 20 : 30 : 45 : 60 41 40 40 35 : 27 39 37 36 : 30 21 35 28 21 16 37 36 : 34 : 25 19 40 39 : 35 : 26 : 20 20 41 40 : 37 : 33 : 28 33 : 28 39 : 39 : 39 : 35 : 26 : 20 20 41 : 40 : 37 : 33 : 28 33 : 28 36 : 36 : 36 : 36 : 36 : 32 : 26 36 : 36 : 36 : 37 : 38 : 38 : 38 : 38 : 38 : 38 : 38	3.5 : 20 : 30 : 45 : 60 · 80 : 	**3.5 : 20 : 30 : 45 : 60 · 80 : 100 411	**3.5 : 20 : 30 : 45 : 60 * 80 : 100 : 150 : 41	**3.5 : 20 : 30 : 45 : 60 * 80 : 100 : 150 : 200 *** 41	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$

Table 49. Hydraulic conductivity of the soil at different suctions.

Danth								0	esorption	Desorption pressure	(E)			
dob	3.5		20		30		45		: 09	80	100	150	200	345
5									cm/hr	r				
0-10	2.02E	10	1.80E	0	1.70E	10		00	2,81E 00	3.59E-01	9.48E-02	1.01E-02	2.04E-03	1,43E-04
10-20	1.45E	. [0	1.18E	0	9.72E	00	4.22E 0	0	8,23E-01	1.46E-01	4.64E-02	4.23E-03	5.96E-04	m
20-30	9.64E	00	9.40E	00	8, 18E	00	3.20E 0	0	7.14E-01	1.50E-01	5.28E-02	6.25E-03	1.44E-03	3.48E-05
30-40	1.84E	5	1,66E	0	1.18E	5	3.22E 0	0	7.87E-01	1.69E-01	5.48E-02	7.94E-03	2.29E-03	3.02E-05
40-50	2,63E	0	2,24E	5	1.35E	0	3.75E O	0	9.70E-01	1.79E-01	5.42E-02	6.50E-03	1.42E-03	4.42E-05
20-60	2.89E	0	2,39E	0	1.72E	0	1.01E 0	_	4.78E 00	1.80E 00	3.15E-01	1.05E-02	1.89E-02	7.90E-05
0/-09	1.97E	0	1.98E	0	2.00E	0	1.31E 0	_	4.16E 00	4.59E-01	1,12E-01	1,03E-01	1.56E-03	5.58E-05
70-80	1.25E (0	1,24E	0	1,23E (0	7.94E 00	0	3.48E 00	3.81E-01	8.21E-02	6.40E-03	9.65E-04	5.65E-05
80-90	2.76E	0	2.53E	0	2,10E	0	1.29E 0	_	2.85E 00	2,63E-01	7.47E-02	3.68E-03	5.87E-04	2,13E-05
+06	2.76E (0	2.59E	0	2,55E	0	1.53E 0	_	3.43E 00	3.09E-01	7.04E-02	4.90E-03	6.46E-04	2.67E-05

Table 50. Quantity-intensity (Q/I) relation of Lakeland fine sand.

0-10		/102 / .067	059 .069	011 .067	007 .066	+.010 .065	+.0	+.055 .083	+.046 .094
19-20	∆K AR	102 .069	059 .069	008 .068	007 .066	+.011 .074	+.011	+.022 .091	+.040 .088
20-30	∆K	070	034	018	067	+.015	+.015	+.055	+.056
	AR	.067	.067	.066	.066	.072	.075	.081	.092
30-40	∆K	102	034	017	013	+.011	+.027	+.046	+.040
	AR	.068	.068	.066	.063	.073	.077	.082	.091
40-50	∆K	102	034	017	005	+.019	+.014	+.046	+.040
	AR	.069	.066	.064	.066	.071	.074	.079	.092
50-60	ΔK	102	025	017	005	+.014	+.014	+.027	+.040
	AR	.069	.067	.066	.067	.076	.074	.082	.070
60-70	∆K AR	102 .068	034 .068	011 .068	007 .069	+.011 .072	+.029 .079	+.040	+.064 .092
70-80	∆K	102	034	011	007	+.014	+.027	+.055	+.000
	AR	.067	.068	.064	.065	.073	.075	.082	.093
80-90	∆K	102	025	008	005	+.002	+.100	+.027	+.040
	AR	.064	.066	.068	.067	.073	.076	.083	.092
90+	△K	102	025	008	007	+.018	+.014	+.027	+.055
	AR	.062	.065	.066	.068	.076	.077	.081	.091

 $[\]frac{1}{2} \text{/Amount of K (me/100 g) released (-) or adsorbed (+)} \\ \text{2} \text{/Activity ratio of equilibrium solution.}$

difficult to present the Q/I relation in graphical form. The data however indicated that the amount of K released or adsorbed by the soil ranged from -0.1 to about +0.05 me/100 g, or about -40 to +20 ppm K in the soil, respectively. This suggests that the amount of K obtained from the soil by equilibrating the soil with NNH $_{\rm L}$ OAc at both pH levels (Table 47) underestimates the amount of K on the exchange complex. The presence of other cations on the exchange complex with greater affinity than K probably reduced adsorption of K by this soil.

The water content at different depths in the column initially and at various times after application of water are given in Tables 51, 52, 53, and 54 for the first, second, third, and fourth water applications, respectively. The K concentration of the soil solution at different depths in the column collected 15 hr after each application of water is shown in Figure 4.

At the start of the experiment, the soil at 90 cm depth was saturated (Table 51), presumably due to the capillary fringe of the water table which was located at 120 cm below the soil surface. There was an increase in water content in the upper portion of the column immediately following the first addition of water which decreased gradually with time following redistribution of water to the lower part of the column. The downward movement of water affected an increase in water content as far as 60 cm depth after 9 hr but this does not necessarily mean that the applied water had reached 60 cm. The applied water may be pushing down the solution initially present in the soil and if a soil layer below has relatively low conductivity, it would result in an accumulation of water above that layer. The second, third, and fourth applications of water resulted in similar water contents in the column prior to each application.

Table 51. Soil water tension (h) and water content (θ) before (t=0) and after the first application of 2.5 cm of water at the rate of 2.5 cm/hr $^{1/2}$

Depth	: t=0 : t=1			: t=1.	. 25	t=	3	: t:	t=4	
рерии	; h	θ	h	9	h	θ	h	θ	h	θ
cm	Cm	%	cm	%	cm	%	cm	%	cm	%
0-10 10-20 20-30 30-40 40-50 50-60 60-70 70-80 80-90 90-100	118 109 92 88 78 68 58 47 37	15 13 15 13 15 19 27 31 35 37	15 97 92 88 78 68 58 47 37 27	41 14 15 13 15 19 27 31 35 37	23 48 92 88 78 68 58 47 37 27	40 28 15 13 15 19 27 31 35 37	47 43 84 87 78 68 58 47 37 27	34 31 15 14 15 19 27 31 35 37	55 44 62 70 75 68 58 47 37 27	29 30 21 16 15 19 27 31 35 37
	t=9		_t=	_t=15		27_	t=/	+5	t=108	
0-10 10-20 20-30 30-40 40-50 50-60 60-70 70-80 80-90 90-100	71 67 69 70 74 66 56 46 37 27	21 19 18 16 15 19 28 32 36 37	82 77 77 73 74 66 56 46 37 27	18 17 16 16 15 19 28 32 36 37	96 88 89 85 78 68 58 47 37 27	16 15 14 15 19 27 31 36 37	106 104 97 88 78 69 59 49 38 28	15 14 14 13 15 18 27 31 35 37	116 118 100 90 80 70 60 50 40	15 13 14 13 14 18 26 30 35 37

 $[\]underline{1}^f$ "t" refers to time in hours after application of water.

Table 52. Soil water tension (h) and water content (θ) before and at different times (in hours) after the second application of 2.5 cm of water at the rate of 2.5 cm/hr.

Depth	Before		t=15		t=24		t=33.5		t=58		t=72	
	h	θ	h	θ	h	θ	h	9	h	θ	h	9
cm	cm	%	cm	%	cm	%	cm	%	cm	%	cm	%
0-10	117	15	81	18	91	17	99	16	112	15	118	15
10-20	118	13	76	17	84	16	93	15	105	14	109	13
20-30	100	14	77	16	82	16	88	15	99	14	101	14
30-40	90	13	77	15	79	15	84	14	90	13	91	13
40-50	80	14	78	15	74	15	77	15	80	14	82	14
50-60	70	18	70	18	66	20	68	19	71	18	72	17
60-70	60	26	59	27	56	28	57	27	61	25	62	24
70-80	50	30	49	30	46	32	48	31	51	30	52	29
80-90	40	35	39	35	36	36	38	35	41	34	42	34
90-100	29	37	30	36	26	37	28	37	31	36	31	36

Table 53. Soil water tension (h) and water content (θ) before and at different times (in hours) after the third application of 2.5 cm of water at the rate of 2.5 cm/hr.

Depth	Before		t:	=15	t=120		
	h	θ	h	θ	h	θ	
cm	cm	%	cm	%	cm	%	
0-10	118	15	88	17	-	_	
10-20	109	13	82	16	111	13	
20-30	100	14	82	16	96	15	
30-40	91	13	79	15	93	13	
40-50	82	14	75	15	84	14	
50-60	72	17	66	19	75	17	
60-70	62	24	59	27	65	22	
70-80	52	30	49	31	55	28	
80-90	42	34	38	35	45	33	
90-100	31	36	28	37	35	36	

Table 54. Soil water tension (h) and water content (0) before and 15 hr after the fourth application of 2.5 cm of water at the rate of 2.5 cm/hr.

Depth	Bef	ore	After 15 hr		
op til	h	θ	h	9	
cm	cm	%	cm	%	
0-10	120	14	78	19	
0-20	111	13	72	18	
0-30	95	14	74	17	
0-40	93	13	83	14	
0-50	84	14	80	15	
0-60	74	17	71	18	
0-70	65	22	61	25	
08-0	55	28	51	30	
0-90	45	33	41	34	
0-100	35	36	31	36	

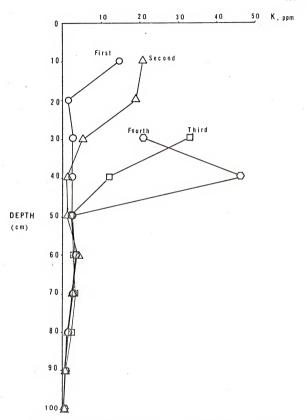


Figure 4. Potassium concentration in soil solution at different depths 15 hr after the first, second, third, and fourth application of 2.5 cm of water at 2.5 cm/hr rate.

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Data on K concentration of the soil solution at 10 and 20 cm depth for the third and fourth applications of water were not obtained because of the difficulty in collecting enough volume of solution for analysis. It is very apparent however, that K moved downward in the soil column as a result of movement of the applied water. The front of the water applied through the fourth application theoretically should reach the 60 cm depth assuming the following: 1) the applied water displaced the water initially present in the column, 2) the initial water content was attained after redistribution of the applied water, 3) there was no buildup in water content at sampling time, and 4) no removal of water occurred (by evaporation for example) except through the bottom of the column when maintaining the hydrostatic pressure at 120 cm below the soil surface where the water table was initially located. Collecting soil solution samples would decrease the value of the theoretical depth. Since the amount of each sample collected in the experiment never exceeded 3 ml, the maximum decrease would be only 10 cm which means that the applied water should then theoretically reach 50 cm after redistribution. However after 15 hr from the fourth application of water, the K concentration of the soil solution at 50 cm depth was just slightly over the initial K concentration while the K concentration at 40 cm depth was highest. This would indicate that K was being held back. Potassium underwent ionic exchange spontaneously during flow of the solution through the column.

Using the second column and after continuous application of 15 cm of water, the concentrations of K, Ca, and Mg in the soil solution and water content at different depths following redistribution of the applied water are given in Table 55. The maximum K concentration in the soil

Table 55. Soil water tension (h), water content (0) and K, Ca, and Mg contents of the soil solution at different depths before and 9.5 hr after 15 cm of continuous application of water.

Depth	Before		After 9.5 hr					
	h	9	h	9	K	Ca	Mg	
Cm	cm	%	cm	%		- ppm		
0-10	105	15	44	36	6.3	5.2	2.0	
10-20	91	15	48	28	7.6	8.0	1.3	
20-30	80	16	53	24	14.2	13.0	2.7	
30-40	70	16	59	20	55.0	27.0	3.1	
40-50	60	20	48	25	212.5	235.0	21.5	
50-60	50	26	41	31	1.0	52.0	10.2	
60-70	40	37	32	38	0.3	30.5	4.3	
70-80	30	34	23	35	0.2	16.8	2.5	
80-90	19	39	14	39	0.2	12.3	1.7	
90-100	9	32	6	32	1.0	9.3	1.1	

solution was at 50 cm depth and the concentration decreased gradually towards the surface of the column. If the rate of water application was sufficient to saturate the soil surface and flow occurred under watersaturated conditions according to the saturated water contents (Table 46). then 40 cm would be the theoretical depth reached with the application of 15 cm of water. In reality, this is not expected since, unless there is a layer below the soil surface with physical properties that may restrict water movement, the surface soil will not saturate with 2.5 cm/hr rate of water application because saturated hydraulic conductivity is 19 cm/hr. In the experiment, the surface soil actually never became saturated since water application rate was much less than the hydraulic conductivity at about -45 cm of water tension of the soil. The data indicate that at the end of the water application period, the highest K concentration in the soil solution was at 50 cm depth. Undoubtedly, flow was taking place under unsaturated conditions and apparently, the highest concentration of K was in the advancing bulk solution. During the redistribution of water to approach the initial water content. K moved with water to lower depths in the column. The tailing of K concentration towards the surface may be due to either incomplete dissolution of KCl or dilution.

Although no data on Ca and Mg concentrations were obtained prior to application of water, it appears that Ca and Mg were being pushed down with the displaced solution. Since no Ca or Mg was applied to the column, this data suggest that movement of Ca and Mg in the column was a result of replacement on the soil exchange sites by K. The high K concentration in the solution in dynamic equilibrium with the soil probably resulted in greater adsorption of K by the soil and consequently, greater replacement of Ca and Mg by mass action. In contrast, the low K concentrations

(90 ppm K) of solutions used in the / / relation study resulted in lower K adsorption.

During miscible displacement of one fluid by another, solute diffusion across the boundary results in the disappearance of the initially sharp boundary between the two fluids. The variation in microscopic velocities of fluid during miscible displacement also causes dispersion of the solute in the displacing fluid. Therefore, solute dispersion primarily results from either molecular diffusion or fluid dispersion or both.

Laboratory procedures for studying solute dispersion under saturated condition usually involve saturating a column of soil with a solution and displacing it with another solution containing the solute being studied. The change in solute concentration (C) of the effluent is plotted against the volume of effluent collected at one end of the column. A breakthrough curve is normally obtained by a plot of the dimensionless ratio C/Co (relative concentration) against the number of pore volumes, V/V_{O} , of effluent collected, where C_{O} is the initial concentration of the solute in the solution prior to displacement, V is the volume of effluent. and V_0 is the effective pore volume. Theoretically, if no fluid mixing occurs (such as in a water pipe), the breakthrough curve will be that of a piston flow, i. e., when one pore volume of effluent has been collected the solute concentration in the effluent will abruptly change to that of the displacing liquid ($C/C_0=1$). Experimental breakthrough curves for porous media, however, usually show a gradual increase followed by a decrease in concentration of the solute in the effluent as the displacing fluid passes through the column. Equations for one-dimensional dispersion of an inert solute (such as C1-) dictate that when one pore volume has

passed through the column, C/C_0 should be equal to 0.5. The position as well as the shape of the breakthrough curve for certain solutions with reference to this point characterizes solute dispersion with regard to adsorption in the soil. Adsorption of the solute causes a delay in the breakthrough, i. e., the curve will appear to the right of the theoretical point $(C/C_0=0.5)$. Desorption of the solute may result in extensive tailing of a breakthrough curve. Repulsion of the solute by the soil causes an advanced appearance of the breakthrough curve.

The experimental breakthrough curves for K and Cl in Lakeland fine sand are shown in Figures 5 to 8. The C1⁻ concentrations in the effluent fractions were based on ${\rm Cl}^{36}$ activity. The curves for K were always to the right of C1 curve. It appears that C1 may have been adsorbed by the soil but to a lesser extent than K. Under saturated conditions (Figures 5 to 7), the effect of increasing flow velocity from 18 to 22 cm/hr was to cause the K breakthrough curve to appear earlier at the faster flow rate than at the lower flow rate. This is probably an effect of pushing the solution through the larger pores faster. Also, the peak of the K breakthrough curve decreased with increased flow velocity while that of Cl did not change. There was no difference in the breakthrough curves between flow velocities of 22 and 69 cm/hr except that the peak of K breakthrough curve was slightly lower at the higher flow velocity than at the lower flow velocity. The breakthrough curves for unsaturated conditions (Figure 8) show an increased delay of K breakthrough with reference to C1 and to the reference point (C/C $_0$ =0.5). It appears that C1 is being repelled by the soil since the C1 breakthrough curve appeared earlier relative to the theoretical point.

Desaturation changes the pattern of flow of the solution through the

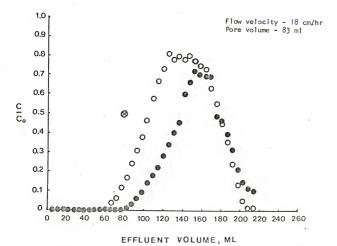


Figure 5. Breakthrough curves of K () and C1 () under saturated condition at flow velocity of 18 cm/hr.

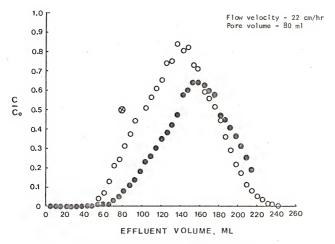


Figure 6. Breakthrough curves of K (♠) and C1 (♠) under saturated condition at flow velocity of 22 cm/hr.

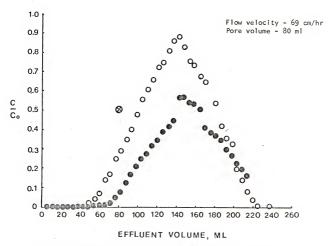
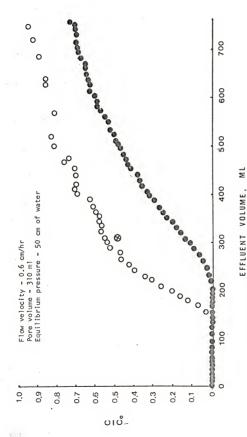


Figure 7. Breakthrough curves of K (●) and C1 (○) under saturated condition at flow velocity of 69 cm/hr.



Breakthrough curves of K () and C1 (O) under unsaturated condition. Figure 8.

soil. The contribution of the larger pores to the flow diminishes as the water content decreases while the smaller pores increase their importance. Consequently, the flow path becomes more tortuous and the proportion of the solution in contact with the soil particles participating in ion exchange increases.

The delay in the breakthrough of K both under saturated and unsaturated conditions can be attributed to adsorption of K on the soil exchange complex. The increased delay for the unsaturated flow was probably due to greater K adsorption than for the saturated flow. The slower velocity of flow under unsaturated conditions allows completion of ion exchange between soil and solution.

SUMMARY AND CONCLUSIONS

Two field experiments were conducted to study K movement from soluble and slowly soluble K fertilizers and its effect on yield and composition of flue-cured tobacco.

In the 1972 experiment, the main treatments consisted of the tobacco-grade fertilizer (4-3.5-10) applied split or once; compounded 4-3.5-10 with K supplied as sulfur-coated $\rm K_2SO_4$ (termed SCK) with 36% coating or with 18% coating, and as uncoated granular $\rm K_2SO_4;$ and resincoated 7-6-17.5 (Osmocote). Subtreatments were three rates equivalent to 450, 900, and 1800 kg/ha of 4-3.5-10. In addition, a treatment with no K (4-3.5-0) and a treatment using Osmocote 7-6-17.5 with all N applied at planting were studied. The main treatments did not significantly affect yields, total alkaloid contents and reducing sugar contents of the leaves. The K contents of the leaves were higher for tobacco-grade 4-3.5-10 applied split, Osmocote 7-6-17.5, and with granular $\mathrm{K}_2\mathrm{SO}_{\underline{L}}$ than the remaining treatments. The K content of the soil samples collected at different depths varied significantly among treatments but the effect of treatments was independent of sampling. The soil samples from plots treated with tobacco-grade 4-3.5-10 applied split and Osmocote 7-6-17.5 contained significantly less K than plots treated with granular $\mathrm{K_2SO}_{\underline{\mu}}$. Data on K uptake could not explain the differences in the K contents of the soil and there was no significant treatment x depth interaction effect. The data indicated greater movement of K from soluble K fertilizers to lower depths than from the slowly soluble K fertilizers. Because of the relatively high rates and low rainfall during the experiment, treatments did not affect yield or

composition of flue-cured tobacco.

In the 1973 experiment, the main treatments consisted of the tobacco-grade 4-3.5-10 applied split, Swift product 0-6.5-25 with N supplied either from IBDU or $\mathrm{NH_4NO_3}$, compounded 4-3.5-10 with K supplied as SCK (26% coating), Osmocote 7-6-17.5, and 4-3.5-0. Subtreatments were three rates similar to the 1972 experiment. The Swift product. SCK, and Osmocote were the three slowly soluble K fertilizers in the study. Yields of flue-cured tobacco varied among fertilizer treatments in two out of five primings as well as the total of the five priming yields. Higher total yields were obtained from the slowly soluble K fertilizers than the soluble one. The K contents of the leaves varied significantly in the first, second, and fifth priming. The slowly soluble K fertilizers, except where IBDU was applied, generally increased K contents of the leaves and K uptake. Where no K was applied, the K contents of the leaves did not appreciably differ from those where Kwas applied. The very wet condition in the 1973 experiment may have moved K laterally and affected the no-K treated plots in the experimental area. Total alkaloid content of the leaves increased with rate while reducing sugar content decreased with rate of fertilizer applied. The effect of treatments on total reducing sugar contents seemed to be more associated with N availability than K. There were differences in the value of flue-cured tobacco but the differences did not indicate any advantage of the slowly soluble K fertilizers over the tobacco-grade fertilizer. The K contents of the soil were lower than those in the 1972 experiment. The K from the slowly soluble K fertilizers did not seem to move down in the soil as quickly as the tobacco-grade fertilizer possibly because they did not dissolve as readily.

Field and laboratory experiments were conducted to study effect of water movement on K movement in the soil. In the field, results showed that intermittent application of water totalling 11.18 cm did not move K from a soluble fertilizer (7-6-17.5) lower than about 15 cm in the profile. With continuous application of 15 cm of water, the K from soluble fertilizer moved down to as far as 75 cm while most K from slowly soluble form of 7-6-17.5 was still in the upper 45-cm depth. The results from laboratory studies suggested that K was lagging behind the water moving downward possibly due to adsorption by the soil. This observation was supported by the data from miscible displacement studies for K and C1⁻ under saturated and unsaturated condition.

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I certify that I have read this study and that in my opinion it conforms to acceptable standard of scholarly presentation and is fully adequate, in scope and quality, as a dissertation for the degree of Doctor of Philosophy.

William K. Robertson, Chairman Professor of Soil Science

I certify that I have read this study and that in my opinion it conforms to acceptable standard of scholarly presentation and is fully adequate, in scope and quality, as a dissertation for the degree of Doctor of Philosophy.

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